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(54) **Novel fibrinogen receptor antagonists.**

(57) A series of non-peptide derivatives that are antagonists of the fibrinogen IIb/IIIa receptor and thus are platelet aggregation compounds useful in the prevention and treatment of diseases caused by thrombus formation.

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BACKGROUND OF THE INVENTION

The present invention provides novel compounds, novel compositions, methods of their use and methods of their manufacture, such compounds being generally pharmacologically useful as anti-platelet aggregation agents in various vascular pathologies. The aforementioned pharmacologic activities are useful in the treatment of mammals. More specifically, the compounds of the present invention act by blocking the platelet receptor site of the protein fibrinogen. Fibrinogen is a glycoprotein that circulates in the blood plasma, and whose platelet receptor site is glycoprotein IIb/IIIa. By blocking the action of fibrinogen at the receptor (glycoprotein IIb/IIIa), the compounds of the present invention interfere with platelet aggregation, which is a cause of many vascular pathologies. At the present time, there is a need in the area of vascular therapeutics for such a fibrinogen receptor blocking agent. By interfering with hemostasis, such therapy would decrease the morbidity and mortality of thrombotic disease.

Hemostasis is the spontaneous process of stopping bleeding from damaged blood vessels. Precapillary vessels contract immediately when cut. Within seconds, thrombocytes, or blood platelets, are bound to the exposed matrix of the injured vessel by a process called platelet adhesion. Platelets also stick to each other in a phenomenon known as platelet aggregation to form a platelet plug. This platelet plug can stop bleeding quickly, but it must be reinforced by the protein fibrin for long-term effectiveness, until the blood vessel tear can be permanently repaired by growth of fibroblasts, which are specialized tissue repair cells.

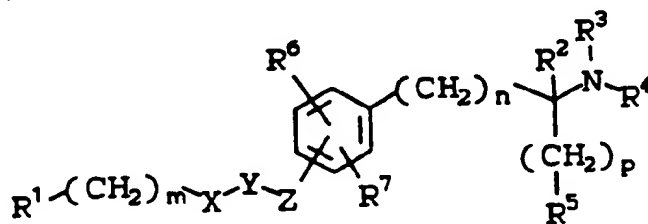
An intravascular thrombus (clot) results from a pathological disturbance of hemostasis. The thrombus can grow to sufficient size to block off arterial blood vessels. Thrombi can also form in areas of stasis or slow blood flow in veins. Venous thrombi can easily detach portions of themselves called emboli that travel through the circulatory system and can result in blockade of other vessels, such as pulmonary arteries. Thus, arterial thrombi cause serious disease by local blockade, whereas venous thrombi do so primarily by distant blockade, or embolization. These diseases include venous thrombosis, thrombophlebitis, arterial embolism, coronary and cerebral arterial thrombosis, myocardial infarction, stroke, cerebral embolism, kidney embolisms and pulmonary embolisms.

There is a need in the area of cardiovascular and cerebrovascular therapeutics for an agent which can be used in the prevention and treatment of thrombi, with minimal side effects, including unwanted prolongation of bleeding in other parts of the circulation while preventing or treating target thrombi. The compounds of the present invention meet this need in the art by providing therapeutic agents for the prevention and treatment of thrombi.

The compounds of the present invention show efficacy as antithrombotic agents by virtue of their ability to block fibrinogen from acting at its platelet receptor site, and thus prevent platelet aggregation.

SUMMARY OF THE INVENTION

The present invention relates to novel compounds having the general structural formula I:

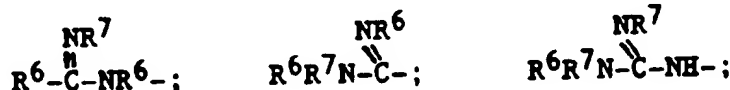


I

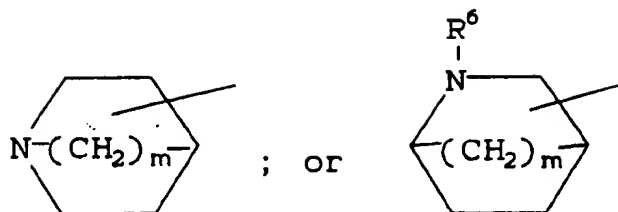
and the pharmaceutically acceptable salts thereof, wherein

R¹ is

a four to eight member heterocyclic ring containing 1, 2, 3 or 4 hetero atoms wherein said heteroatoms are N, O or S and wherein said heterocyclic ring is optionally substituted at any atom by H, R⁶ or R⁷;



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NR⁶R⁷ wherein R⁶ and R⁷ are independently hydrogen,
C₁₋₁₀ alkoxycarbonyl or unsubstituted or substituted C₁₋₁₀ alkyl and cycloalkyl wherein said substituents are

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C₁₋₁₀ alkoxy,

C₁₋₁₀ alkoxyalkyl,

C₁₋₁₀ alkoxyalkyloxy,

C₁₋₁₀ alkoxycarbonyl,

C₁₋₁₀ alkylcarbonyl,

C₀₋₈ alkylaminocarbonyl,

C₁₋₁₀ aralkylcarbonyl,

C₁₋₁₀ alkylthiocarbonyl,

C₄₋₁₀ aralkylthiocarbonyl, thiocarbonyl,

C₁₋₁₀ alkoxythiocarbonyl, aryl,

5 to 6 membered saturated heterocyclic rings of 1, 2, 3 or 4 hetero atoms wherein said hetero atoms are taken from the group consisting of N, O and S,

C₁₋₄ alkanoylamino,

C₁₋₈ alkoxycarbonyl-C₀₋₈ alkylamino,

C₁₋₁₀ alkylsulfonylamino,

C₄₋₁₀ aralkylsulfonylamino,

C₄₋₁₀ aralkyl,

C₁₋₁₀ alkaryl,

C₁₋₁₀ alkylthio,

C₄₋₁₀ aralkylthio,

C₁₋₁₀ alkylsulfinyl,

C₄₋₁₀ aralkylsulfinyl,

C₁₋₁₀ alkylsulfonyl,

C₄₋₁₀ aralkylsulfonyl, aminosulfonyl,

C₁₋₁₀ alkylaminosulfonyl,

C₄₋₁₀ aralkylsulfonylamino,

oxo,

thio,

unsubstituted or mono- or di-substituted 1-ethenyl, 2-ethenyl or 3-propenyl wherein said substituents are selected from the group consisting of hydrogen, C₁₋₁₀ alkyl and C₇₋₁₀

aralkyl,

carboxy,

hydroxy,

amino,

C₁₋₈ alkylamino,

C₁₋₈ dialkylamino,

halogen, where halogen is defined as C1, F, Br, or I,

nitro, or

cyano,

and further wherein said N can additionally be substituted to form a quaternary ammonium ion wherein said substituent is as previously defined for R⁶ and R⁷;

R² and R³ are independently
hydrogen,

aryl or

unsubstituted or substituted C₀₋₁₀ alkyl or cycloalkyl wherein said substituent is

C₁₋₁₀ alkoxyalkyl,

aryl,

a 4 to 8 membered heterocyclic ring containing 1, 2, 3 or 4 hetero atoms, wherein said heteroatoms are taken from the group consisting of N, O and S,

C₄₋₁₀ aralkyl,

C₁₋₁₀ alkaryl, carboxy,

C₁₋₁₀ alkylcarbonyl,

C₁₋₁₀ alkylthiocarbonyl,

C₄₋₁₀ aralkylcarbonyl,

C₄₋₁₀ aralkylthiocarbonyl,

C₁₋₆ alkoxy carbonyl,

C₄₋₁₀ aralkoxy carbonyl,

C₁₋₆ alkoxy,

C₄₋₁₀ aralkoxy,

C₁₋₆ alkylamino,

C₁₋₁₂ dialkylamino,

C₁₋₆ alkanoylamino,

C₄₋₁₂ aralkanoylamino,

C₄₋₁₀ aralkylamino;

R⁴ is

hydrogen,

aryl,

C₁₋₁₀ alkyl or cycloalkyl

C₄₋₁₀ aralkyl,

arylcarbonyl, aminocarbonyl,

C₁₋₁₀ alkylcarbonyl, C₁₋₆ alkylaminocarbonyl,

C₁₋₁₀ alkylthiocarbonyl, C₁₋₆ dialkylamino-carbonyl,

C₁₋₁₀ alkoxythiocarbonyl, arylC₁₋₆ alkylamino-carbonyl,

C₁₋₁₀ alkoxy carbonyl,

C₄₋₁₀ aralkylcarbonyl,

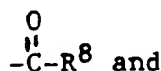
C₄₋₁₀ aralkoxy carbonyl,

C₁₋₁₀ carboxyalkyl and

further wherein any of the substituents for R⁴ may be substituted by one or more substituents selected from the group as defined for R⁶, or an L- or D-amino acid joined by an amide linkage;

R⁵ is

a four to eight membered saturated or unsaturated heterocyclic ring containing 1, 2, 3 or 4 hetero atoms wherein said hetero atoms are N, O, or S or,



wherein R⁸ is

hydroxy,

C₁₋₁₀ alkyloxy,

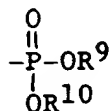
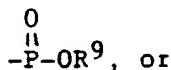
C₁₋₁₀ alkaryloxy,

C₄₋₁₀ aralkyloxy,

C₄₋₁₀ aralkylcarbonyloxy,

C₁₋₁₀ alkoxyalkyloxy,

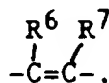
C_{1-10} alkoxyalkylcarbonyloxy,
 C_{1-10} alkoxyalkylcarbonyloxyalkyl,
 C_{1-10} alkylcarbonyloxyalkyloxy,
 an L- or D-amino acid joined by an amide linkage, and wherein the carboxylic acid moiety of said amino
 acid is as the free acid or is esterified by C_{1-6} alkyl.



wherein R^9 and R^{10} are selected from the group consisting of
 hydrogen, C_{1-10} alkyl and C_{4-10} aralkyl;

X and Y are optional substituents that, when present, are

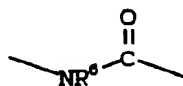
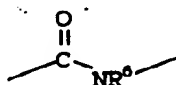
NR^6 ,
 O ,
 S ,
 SO ,
 SO_2 ,



$-\text{C}=\text{C}-$,
 oxo,
 aryl,
 thiono,

unsubstituted or substituted C_{1-15} alkyl or cycloalkyl wherein said substituents are independently R^6 and

R^7 ,



$-\text{NR}^6-\text{SO}_2-$, $-\text{SO}_2-\text{NR}^6-$, or

a 4- to 8- membered heterocyclic ring containing 1, 2, 3, or 4 heteroatoms wherein said atoms are N, O, or S
 and wherein said ring is independently substituted at any atom with R^6 ;

Z is an optional substituent that, when present, is independently chosen as defined by X and Y;

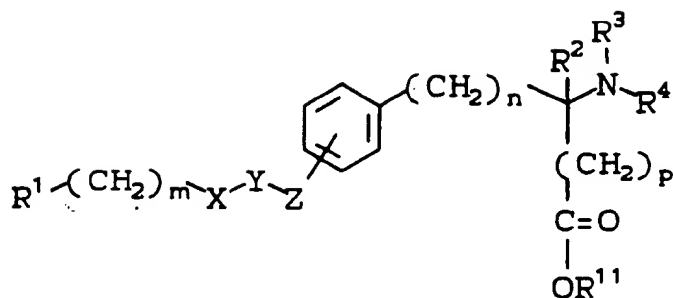
m is an integer of from zero to ten;

n is an integer of from zero to ten; and

p is an integer of from zero to three.

A preferred group of compounds of the present invention are those defined for general structural formula

Il as:



II

wherein

R^1 is

a five to six membered heterocyclic ring wherein said heteroatoms are N, O, or S, and wherein said heterocyclic ring is optionally substituted by hydrogen, C₁₋₅ alkyl; or

NR⁶R⁷ wherein R⁶ and R⁷ are independently hydrogen, unsubstituted or substituted C₁₋₁₀ alkyl or C₄₋₁₀ aralkyl wherein said substituents are chosen from

C₁₋₁₀ alkoxy carbonyl,

anyt,

C₀₋₅ dialkylamino C₁₋₁₀ alkyl, and

C₄₋₁₀ aralkyl,

and further wherein said N can additionally be substituted to form a quaternary ammonium ion;

R² and R³ are hydrogen, C₁₋₄ alkyl or C₄₋₁₀ aralkyl;

R4 is

 H_1

C₁₋₁₀ alkyl,

C₄₋₁₀ aralkyl,

arylcarbonyl,

C₁₋₁₀ alkylcarbonyl,

C₁₋₁₀ alkoxy carbonyl,

C₄₋₁₀ aralkylcarbonyl, or

C₄₋₁₀ aralkoxycarbonyl,

wherein R⁴ is unsubstituted or substituted with R⁶ as previously defined;

R¹¹ is

hydrogen or

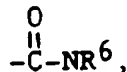
C₁₋₁₀ alkyl;

X and Y are independently

0, S, SO, SO₂,



aryl,

$$-\text{CH}=\text{CH}-,$$




5 $-\text{SO}_2\text{NR}^6$, $-\text{NR}^6\text{SO}_2$, or a 5- or 6- membered heterocyclic ring containing 1 or 2 heteroatoms, wherein said atoms are N, O or S, unsubstituted or substituted C_{1-15} straight, branched, or cyclic alkyl wherein said substituent is oxo, hydroxy C_{1-4} alkyloxy, or C_{4-10} arylalkyl;

Z is an optional substituent that, when present, is

0, SO_2 , $-\text{NR}^6\text{CO}$, $-\text{CONR}^6$,



or

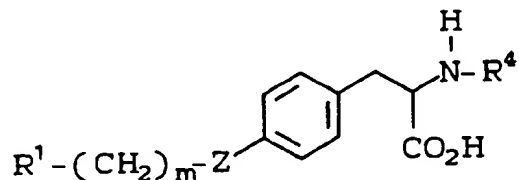
C_{1-10} straight or branched alkyl;

m is an integer of from zero to eight;

n is an integer of from zero to two; and

20 p is an integer of from zero to two.

A more preferred group of compounds of the present invention are those defined for the general structural formula III as



wherein

R^1 is

a five or 6-membered heterocyclic ring wherein said heteroatom is N and wherein said heterocyclic ring is optionally substituted by hydrogen or C_{1-6} alkyl, or

NR^6R^7 wherein R^6 and R^7 are independently hydrogen, C_{1-10} alkyl or C_{4-10} arylalkyl;

R^4 is

arylcarbonyl,

C_{1-10} alkylcarbonyl,

C_{1-10} alkoxy carbonyl,

C_{4-10} aralkylcarbonyl,

C_{4-10} aralkoxy carbonyl wherein R^4 is unsubstituted or substituted with R^6 as previously defined;

Z is chosen from:

0, $-\text{NR}^6\text{CO}$, $-\text{CONR}^6$, or CH_2 ; and

m is an integer of from one to six

DETAILED DESCRIPTION OF THE INVENTION

The term "pharmaceutically acceptable salts" shall mean non-toxic salts of the compounds of this invention which are generally prepared by reacting the free base with a suitable organic or inorganic acid. Representative salts include the following salts:

Acetate

Benzenesulfonate

Benzoate

Bicarbonat

Bisulfate

Bitartrate

	Borate
	Bromide
	Calcium Edetate
	Camsylate
5	Carbonate
	Chloride
	Clavulanate
	Citrate
	Dihydrochloride
10	Edetate
	Edisylate
	Estolate
	Esylate
	Fumarate
15	Glucaptate
	Gluconate
	Glutamate
	Glycolylarsanilate
	Hexylresorcinate
20	Hydrabamine
	Hydrobromide
	Hydrochloride
	Hydroxynaphthoate
	Iodide
25	Isothionate
	Lactate
	Lactobionate
	Laurate
	Malate
30	Maleate
	Mandelate
	Mesylate
	Methylbromide
	Methylnitrate
35	Methylsulfate
	Mucate
	Napsylate
	Nitrate
	Oleate
40	Oxalate
	Pamaote
	Palmitate
	Pantothenate
	Phosphate/diphosphate
45	Polygalacturonate
	Salicylate
	Stearate
	Subacetate
	Succinate
50	Tannate
	Tartrate
	Teoclate
	Tosylate
	Triethiodide
55	Valerate

The term "pharmacologically effective amount" shall mean that amount of a drug or pharmaceutical agent that will elicit the biological or medical response of a tissue, system or animal that is being sought by a researcher or clinician. The term "anti-coagulant" shall include heparin, and warfarin. The term "thrombolytic agent" shall

include streptokinase and tissue plasminogen activator. The term "platelet anti-aggregation agent" shall include aspirin and dipyridimole.

The term "aryl" shall mean a mono- or polycyclic system composed of 5- and 6- membered aromatic rings containing 0, 1, 2, 3 or 4 heteroatoms chosen from N, O or S and either unsubstituted or substituted with R⁶.

5 The term "alkyl" shall mean straight or branched chain alkane, alkene or alkyne.

The term "alkoxy" shall be taken to include an alkyl portion where alkyl is as defined above.

The terms "aralkyl" and "alkaryl" shall be taken to include an alkyl portion where alkyl is as defined above and to include an aryl portion where aryl is as defined above.

The term "halogen" shall include fluorine, chlorine, iodine and bromine.

10 The term "oxo" shall mean the radical =O.

The term "thio" shall mean the radical =S.

In the schemes and examples below, various reagent symbols have the following meanings:

BOC: t-butyloxycarbonyl.
 Pd-C: Palladium on activated carbon catalyst.
 15 DMF: Dimethylformamide.
 DMSO: Dimethylsulfoxide.
 CBZ: Carbobenzyloxy
 CH₂Cl₂: methylene chloride
 CHCl₃: chloroform
 20 EtOH: ethanol
 MeOH: Methanol
 EtOAc: ethylacetate
 HOAc: acetic acid
 BOP: Benzotriazol-1-yloxytris(dimethylamino)-phosphonium, hexafluororophosphate

25 The compounds of the present invention can be administered in such oral dosage forms as tablets, capsules (each of which includes sustained release or timed release formulations), pills, powders, granules, elixers, tinctures, suspensions, syrups and emulsions. Likewise, they may also be administered in intravenous (bolus or infusion), intraperitoneal, subcutaneous or intramuscular form, all using forms well known to those of ordinary skill in the pharmaceutical arts. An effective but non-toxic amount of the compound desired can be employed as an anti-aggregation agent.

30 Compounds of the invention may be administered to patients where prevention of thrombosis by inhibiting binding of fibrinogen to the platelet membrane glycoprotein complex IIb/IIIa receptor is desired. They are useful in surgery on peripheral arteries (arterial grafts, carotid endarterectomy) and in cardiovascular surgery where manipulation of arteries and organs, and/or the interaction of platelets with artificial surfaces, leads to platelet aggregation and consumption. The aggregated platelets may form thrombi and thromboemboli. They may be administered to these surgical patients to prevent the formation of thrombi and thromboemboli.

35 Extracorporeal circulation is routinely used for cardiovascular surgery in order to oxygenate blood. Platelets adhere to surfaces of the extracorporeal circuit. Adhesion is dependent on the interaction between GPIIb/IIIa on the platelet membranes and fibrinogen adsorbed to the surface of the circuit. (Glusko et al., Amer. J. Physiol., 1987, 252:H, pp 615-621). Platelets released from artificial surfaces show impaired hemostatic function. Compounds of the invention may be administered to prevent adhesion.

40 Other applications of these compounds include prevention of platelet thrombosis, thromboembolism and reocclusion during and after thrombolytic therapy and prevention of platelet thrombosis, thromboembolism and reocclusion after angioplasty of coronary and other arteries and after coronary artery bypass procedures. They may also be used to prevent myocardial infarction.

45 The dosage regimen utilizing the compounds of the present invention is selected in accordance with a variety of factors including type, species, age, weight, sex and medical condition of the patient; the severity of the condition to be treated; the route of administration; the renal and hepatic function of the patient; and the particular compound or salt thereof employed. An ordinarily skilled physician or veterinarian can readily determine and prescribe the effective amount of the drug required to prevent, counter or arrest the progress of the condition.

50 Oral dosages of the present invention, when used for the indicated effects, will range between about 0.01 mg per kg of body weight per day (mg/kg/day) to about 100 mg/kg/day and preferably 1.0-100 mg/kg/day and most preferably 1.0 to 20 mg/kg/day. Intravenously, the most preferred doses will range from about 1 to about 55 10 mg/kg/minute during a constant rate infusion. Advantageously, compounds of the present invention may be administered in a single daily dose, or the total daily dosage may be administered in divided doses of two, three or four times daily. Furthermore, preferred compounds for the present invention can be administered in intranasal form via topical use of suitable intranasal vehicles, or via transdermal routes, using those forms of

transdermal skin patches well known to those of ordinary skill in that art. To be administered in the form of a transdermal delivery system, the dosage administration will, of course, be continuous rather than intermittent throughout the dosage regimen.

5 In the methods of the present invention, the compounds herein described in detail can form the active ingredient, and are typically administered in admixture with suitable pharmaceutical diluents, excipients or carriers (collectively referred to herein as 'carrier' materials) suitably selected with respect to the intended form of administration, that is, oral tablets, capsules, elixirs, syrups and the like, and consistent with conventional pharmaceutical practices.

10 For instance, for oral administration in the form of a tablet or capsule, the active drug component can be combined with an oral, non-toxic, pharmaceutically acceptable, inert carrier such as lactose, starch, sucrose, glucose, methyl cellulose, magnesium stearate, dicalcium phosphate, calcium sulfate, mannitol, sorbitol and the like; for oral administration in liquid form, the oral drug components can be combined with any oral, non-toxic, pharmaceutically acceptable inert carrier such as ethanol, glycerol, water and the like. Moreover, when desired or necessary, suitable binders, lubricants, disintegrating agents and coloring agents can also be incorporated
15 into the mixture. Suitable binders include starch, gelatin, natural sugars such as glucose or beta-lactose, corn sweeteners, natural and synthetic gums such as acacia, tragacanth or sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes and the like. Lubricants used in these dosage forms include sodium oleate, sodium stearate, magnesium stearate, sodium benzoate, sodium acetate, sodium chloride and the like. Disintegrators include, without limitation, starch, methyl cellulose, agar, bentonite, xanthan gum and the like.

20 The compounds of the present invention can also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine or phosphatidylcholines.

Compounds of the present invention may also be delivered by the use of monoclonal antibodies as individual carriers to which the compound molecules are coupled. The compounds of the present invention may also
25 be coupled with soluble polymers as targetable drug carriers. Such polymers can include polyvinylpyrrolidone, pyran copolymer, polyhydroxypropylmethacrylamide-phenol, polyhydroxyethylaspartamidephenol, or polyethyleneoxide-polylysine substituted with palmitoyl residues. Furthermore, the compounds of the present invention may be coupled to a class of biodegradable polymers useful in achieving controlled release of a drug, for example, polylactic acid, polyglycolic acid, copolymers of polylactic and polyglycolic acid, polyepsilon caprolactone, polyhydroxy butyric acid, polyorthoesters, polyacetals, polydihydropyrans, polycyanoacrylates and crosslinked or amphipathic block copolymers of hydrogels.

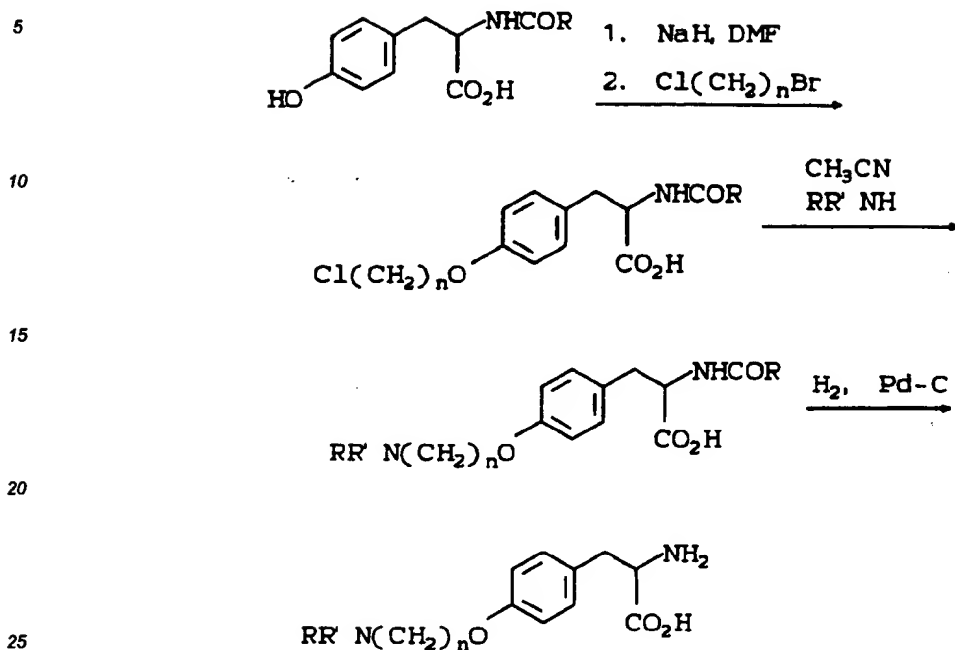
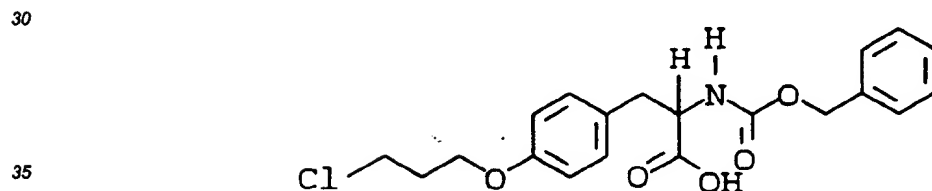
30 The compounds of the present invention can also be co-administered with suitable anti-coagulant agents or thrombolytic agents such as plasminogen activators or streptokinase to achieve synergistic effects in the treatment of various vascular pathologies. They may also be combined with heparin, aspirin or warfarin.

35 The novel compounds of the present invention were prepared according to the procedure of the following schemes and examples, using appropriate materials and are further exemplified by the following specific examples. The most preferred compounds of the invention are any or all of those specifically set forth in these examples. These compounds are not, however, to be construed as forming the only genus that is considered as the invention, and any combination of the compounds or their moieties may itself form a genus. The following examples further illustrate details for the preparation of the compounds of the present invention. Those skilled in the
40 art will readily understand that known variations of the conditions and processes of the following preparative procedures can be used to prepare these compounds. All temperatures are degrees Celsius unless otherwise noted.

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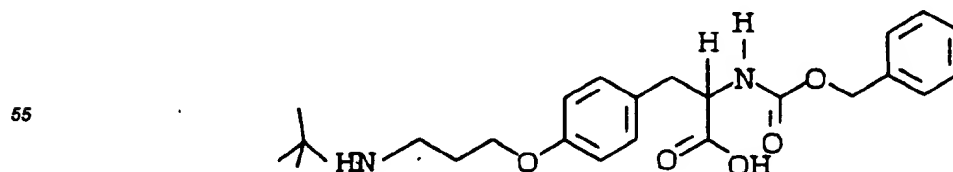
SCHEME 1EXAMPLE 12-S-(N-Benzyloxycarbonylamino)-3-[4-(3-chloropropoxy)phenyl]propionic acid (1-1)

40 N-CBZ-tyrosine (3 g, 9.9 mmole) (from Bachem Chemical Supply, California), was dissolved in DMF and treated with NaH (50% dispersion in oil, 0.95 g, 19.8 mmole) for 1 hour, then 1,3-bromochloropropane (1 ml, 9.9 mmole) was added and the reaction stirred for 16 hours. The DMF was removed in vacuo and the residue dissolved in water, acidified to pH 3, and extracted with ethyl acetate. The ethyl acetate layer was dried with MgSO_4 , filtered and evaporated. Column chromatography (SiO_2 , 97:3:1 $\text{CHCl}_3/\text{CH}_3\text{OH}/\text{HOAc}$) yielded 2.42 g

45 of product as a yellow oil.

RF = 0.3 in 97:3:1 $\text{CHCl}_3/\text{CH}_3\text{OH}/\text{HOAc}$ ninhydrin stain

300 MHz ^1H NMR (CDCl_3) δ 7.3 (bs, 5H), 7.03 (d, $J = 8.3$, 2H), 6.8 (d, $J = 8.3$, 2H), 5.2 (d, $J = 8\text{Hz}$, 1H), 5.05 (bs, 2H), 4.65 (m, 1H), 4.05 (t, $J = 5.7\text{ Hz}$, 2H), 3.73 (t, $J = 6.3\text{ Hz}$, 2H), 3.1 (m, 2H), 2.2 (m, 2H).

EXAMPLE 2

2-S-(Benzyloxycarbonylamino)-[4-(3-t-butylaminopropoxy)phenyl]propionic acid (1-2)

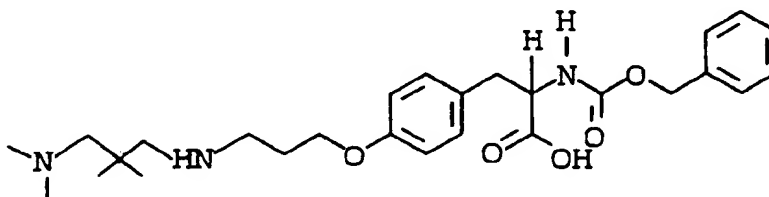
Compound 1-1 (0.4 g, 1.1 mmole) was refluxed in t-butylamine (20 ml) and acetonitrile (20 mL) for three days. The reaction was evaporated to dryness, the residue dissolved in water, and extracted with ether. The aqueous layer was then acidified to pH 4-5 and a precipitate formed. The solid was collected and dried to yield 70 mg of product.

R_f = 0.8 in 9:1 EtOH/NH₄OH, ninhydrin stain.

300 MHz ¹H NMR (D₂O + NaOH) δ 7.4 (bs, 2H), 7.2 (bs, 4H), 6.85 (d, J = 8.55, 2H), 5.2 (d, J = 12.8 Hz, 1H), 5.0 (d, J = 12.8 Hz, 1H), 4.3 (dd, J = 4.0, 9.6 Hz, 1H), 4.0 (bs, 2H), 3.2 (dd, J = 4.0, 13.6 Hz, 1H), 2.8 (dd, J = 9.6 Hz, 13.6 Hz, 1H), 2.65 (t, J = 7.3 Hz, 2H), 1.8 (m, 2H), 1.09 (s, 9H), mass spec (FAB) m/e = 429 (m + 1)
C, H, N analysis C₂₄H₃₂N₂O₆ 0.65 H₂O

MW = 440.244 Calculated C = 65.47, H = 7.62, N = 6.36

Found C = 65.52, H = 7.54, N = 6.27

EXAMPLE 32-S-(N-Benzyloxycarbonylamino)-3-[4-(N,N,2,2-tetramethylpropanediamino)propoxyphenyl]propionic acid (1-3)

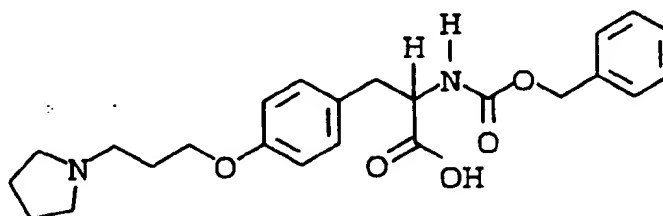
Treatment of compound 1-1 with excess N,N,2,2-tetramethyl-1,3-propanediamine by refluxing in acetonitrile for three days, and followed by an aqueous workup provided crude 1-3. This was chromatographed on silica gel eluting with 9:1:1 EtOH/H₂O/ NH₄OH to provide pure 1-3 (R_f = 0.37 ninhydrin stain). 300 MHz ¹H NMR (D₂O) δ 7.5 (bs, 3H), 7.4 (bs, 2H), 7.33 (d, J = 8.3Hz, 2H), 7.0 (d, J = 8.3Hz, 2H), 5.20 (d, J = 10Hz, 1H), 5.10 (d, J = 10Hz, 1H), 4.25 (m, 1H), 4.25 (t, J = 5.6Hz, 2H), 3.4 (t, J = 7.8Hz, 2H), 3.4 (s, 2H), 3.25-2.95 (m, 2H), 3.22 (s, 2H), 3.1 (s, 6H), 2.35 (m, 2H), 1.38 (s, 6H).

MW = 759.28

C, H, N analysis for C₂₇H₃₉N₂O₅·2.4 C₂HF₃O₂

Calcd: C, 50.30; H, 5.50; N, 5.53.

Found: C, 50.35; H, 5.43; N, 5.56.

EXAMPLE 42-S-(N-Benzyloxycarbonylamino)-3-[4-(3-N-pyrrolidinylpropoxy)phenyl]propionic acid (1-4)

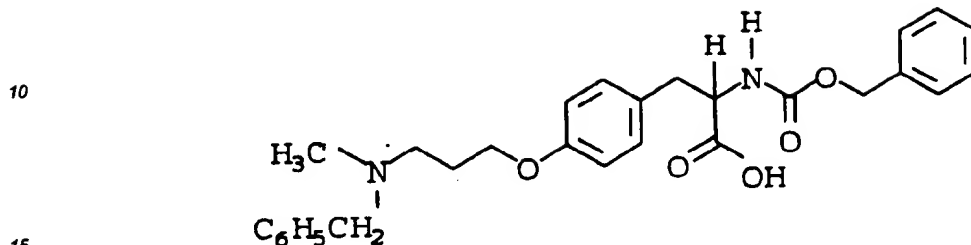
Treatment of compound 1-1 with excess pyrrolidine in CH₃CN at reflux for three days provided crude 1-4. This was purified by flash chromatography on silica gel eluting with 9:1:1 EtOH/H₂O/NH₄OH to give pure 1-4 (R_f = 0.81, ninhydrin stain) in 36% yield. 300 MHz ¹H NMR (CDCl₃) δ 7.3 (bs, 5H), 7.0 (d, J = 8.1Hz, 2H), 6.7 (d, J = 8.1Hz, 2H), 5.5 (d, J = 7.4Hz, 1H), 5.0 (bs, 2H), 4.5 (m, 1H), 3.8 (m, 2H), 3.75 (bs, 1H), 3.4 (m, 2H), 3.18 (t, J = 8.5Hz, 2H), 3.1 (bs, 2H), 2.8 (bs, 1H), 2.2-1.8 (m, 6H).

C, H, N analysis $C_{24}H_{30}N_2O_6 \cdot 0.25 CH_2Cl_2$

Calcd: C, 65.05; H, 6.87; N, 6.26.

Found: C, 65.28; H, 6.78; N, 6.27.

5 EXAMPLE 5



2-S-(N-Benzoyloxycarbonylamino)-[4-(3-N-methyl-N-benzylaminopropoxy)phenyl]propionic acid (1-5)

Treatment of 1-1 with excess N-methyl benzylamine in acetonitrile at reflux for three days afforded crude 1-5. The solvent was removed on a rotary evaporator and the residue was dissolved in water and extracted with 3 x 75 mL portions of ether. The product separated out as an oil which was collected and concentrated to give 1-5 (70 mg) as a foam. 300 MHz 1H NMR ($CDCl_3/CD_3OD$) δ 7.4 (m, 10H), 7.0 (d, $J = 8.5$ Hz, 2H), 6.6 (d, $J = 8.5$ Hz, 2H), 5.0 (bs, 2H), 4.5 (m, 1H), 4.2 (bs, 2H), 3.88 (t, $J = 5.3$ Hz, 2H), 3.1-2.8 (m, 4H), 2.69 (s, 3H), 2.2 (bs, 2H).

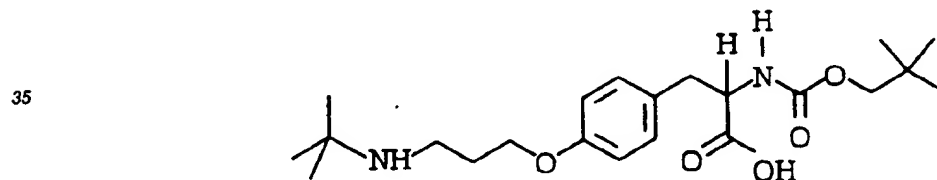
C, H, N analysis $C_{28}H_{32}N_2O_6 \cdot 0.8 CH_2Cl_2 \cdot 0.25 EtOAc$

Calcd: C, 63.17; H, 6.33; N, 4.94.

Found: C, 63.16; H, 6.40; N, 5.04.

MW = 548.771

30 EXAMPLE 6



40 2-S-(N-(t-Butyloxycarbonylamino)-[4-(3-N-t-butylamino propyl-oxy)phenyl]propionic acid (1-6)

Treatment of N-BOC-L-tyrosine with sodium hydride in DMF followed by 1,3-bromochloropropane provided the N-BOC analog of 1-1. This was treated with an excess of t-butylamine in refluxing acetonitrile for two days to provide crude 1-6 after aqueous workup and extraction. Pure 1-6 was prepared by preparative reverse phase HPLC.

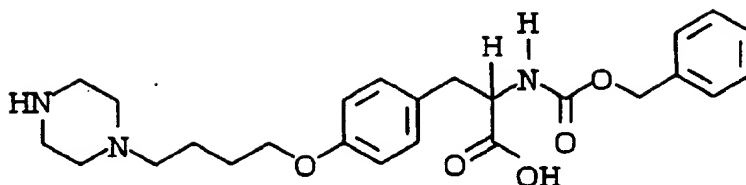
300 MHz 1H NMR (CD_3OD) δ 7.17 (d, $J = 8.5$ Hz, 2H), 6.85 (d, $J = 8.5$ Hz, 2H), 4.28 (dd, $J = 4.8, 9.1$ Hz, 1H), 4.1 (t, $J = 5.9$ Hz, 2H), 3.2 (t, $J = 7.7$ Hz, 2H), 3.1 (dd, $J = 4.8, 13.3$ Hz, 1H), 2.8 (dd, $J = 9.1, 13.3$ Hz, 1H), 2.15 (m, 2H), 1.38 (s, 18H).

C, H, N analysis $C_{21}H_{34}N_2O_7 \cdot 1.05 C_2HF_3O_2$

MW = 514.243

Calcd: C, 53.95; H, 6.87; N, 5.45.

Found: C, 54.01; H, 6.97; N, 5.51.

EXAMPLE 72-S-(N-(Benzyloxycarbonylamino)-3-[4-(4-piperazinyl)butyloxyphenyl]propionic acid (1-7)

Treatment of N-CBZ-L-tyrosine with sodium hydride in DMF followed by 1,4-dibromobutane, as described for the preparation of 1-1, provided the corresponding butyl analog. Treatment of this with 1,4-piperazine in refluxing acetonitrile for three days gave crude 1-7 as a precipitate from the reaction mixture. Reverse phase HPLC purification gave pure 1-7.

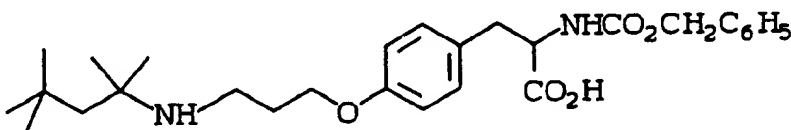
300 MHz ^1H NMR (CD_3OD) δ 7.3 (m, 5H), 7.23 (d, 2H), 6.83 (d, 2H), 5.0 (bs, 2H), 4.35 (dd, 1H), 4.0 (t, 2H), 3.6 (bs, 8H), 3.1 (dd, 1H), 2.85 (dd, 1H), 2.00-1.8 (m, 4H).

C, H, N analysis $\text{C}_{26}\text{H}_{35}\text{N}_3\text{O}_5 \cdot 1.2 \text{H}_2\text{O}$

MW = 491.206

Calcd: C, 63.57; H, 7.67; N, 8.56.

Found: C, 63.33; H, 7.28; N, 8.55.

EXAMPLE 7(a)

1-8

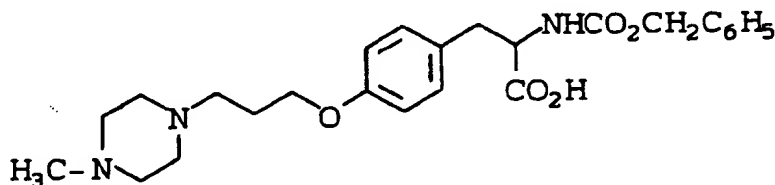
2-S-(N-(Benzyloxycarbonylamino)-3-[4-(1,1,4,4-tetramethylbutylamino)propyloxyphenyl]propionic acid (1-8)

Treatment of 1-1 with 1,1,4,4-tetramethylbutylamine, as described for compound 1-2, gave 1-8 as the TFA salt. ^1H NMR (300 MHz CD_3OD) δ 7.35 (5H, m), 7.18 (2H, d), 6.85 (1H, d), 5.00 (2H, s), 4.35 (1H, dd), 4.10 (2H, t), 3.1 (2H, t), 3.15 (1H, dd), 2.50 (1H, dd), 2.1 (2H, m), 1.70 (2H, s), 1.5 (6H, s), 1.10 (9H, s).

Analysis for $\text{C}_{28}\text{H}_{40}\text{N}_2\text{O}_5 \cdot 0.9 \text{C}_2\text{HF}_3\text{O}_2$

Calcd: C, 60.94; H, 7.02; N, 4.77.

Found: C, 60.85; H, 7.01; N, 4.69.

EXAMPLE 7(b)

1-9

2-S-(N-(Benzyloxycarbonyl)-3-[4-(4-methylpiperazin-1-yl)propyloxyphenyl]propanoic acid (1-9)

Treatment of 1-1 with N-methylpiperazine as described for 1-2 gave crude 1-9. This was purified by column chromatography on silica gel eluting with 9:1:1 $\text{C}_2\text{H}_5\text{OH}/\text{H}_2\text{O}/\text{NH}_4\text{OH}$ to give pure 1-9 as the TFA salt.

^1H NMR (300 MHz D_2O) δ 7.5 (3H, m), 7.4 (2H, d), 7.0 (2H, d), 5.18 (1H, d), 5.05 (1H, d), 4.5 (1H, m), 4.2 (2H,

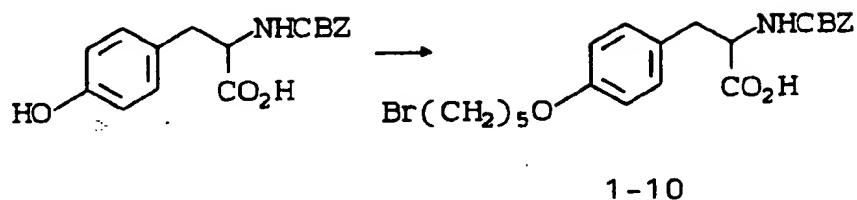
t), 3.8 (8H, s), 3.6 (2H, t), 3.3 (1H, m), 3.1 (3H, s), 3.0 (1H, m), 2.4 (2H, m).

Analysis for $C_{25}H_{33}N_3O_5 \cdot 2.3 C_2HF_3O_2$

Calcd: C, 49.52; H, 4.96; N, 5.85.

Found: C, 49.42; H, 4.98; N, 6.01.

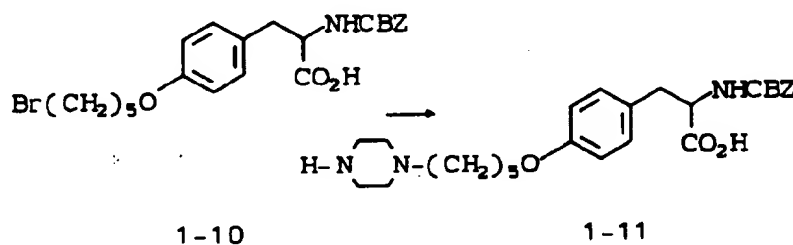
EXAMPLE 7(c)



2-(N-(Benzyloxycarbonylamino)-3-[4-(5-bromopentyloxy)phenyl]propionic acid (1-10)

N-CBZ-L-tyrosine (2.06 g, 5.86 mmole) was treated with NaH (0.58 g, 12.08 mmole) and 1,5-dibromopentane (0.8 ml, 5.87 mmole) as described for 1-1 in Example 1. The crude product was dissolved in methanol and after stirring with silica gel for 0.5 hour, the solvent was removed. This was dry packed and eluted on a flash column with $CHCl_3$ and then with 97:3:0.3 $CHCl_3/CH_3OH/HOAc$ to give pure 1-10 (0.66 g). 1H NMR (300 MHz, CD_3OD) δ 1.50-1.65 (2H, m), 1.63-1.95 (4H, m), 3.10 (2H, m), 3.45 (1H, t), 3.92 (2H, m), 4.40 (1H, m), 6.80 (2H, d), 7.10 (2H, d), 7.28 (5H, m).

EXAMPLE 7(d)



2-S-(N-(Benzyloxycarbonylamino)-3-[4-(4-piperazin-1-yl)-pentyloxyphenyl]propionic acid (1-11)

1-10 (0.658 g, 1.42 mmole), was dissolved in 30 mL CH_3CN and 1,4-piperazine (1.22 g, 14.16 mmole) was added. This solution was stirred at room temperature for 4 days. The solvent was then removed and the residue was dry packed on a silica gel column and eluted with 18:1:1 $C_2H_5OH/H_2O/NH_4OH$ to give pure 1-11 (34 mg) as a white solid.

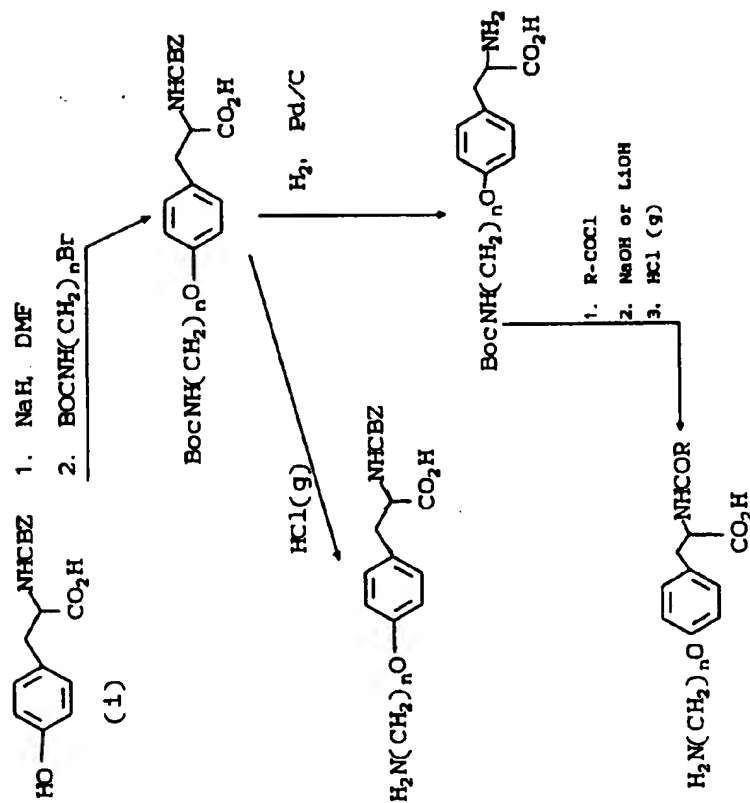
1H NMR (300 MHz, CD_3OD) δ 1.52 (4H, m), 1.77 (2H, m), 2.40 (2H, t), 2.59 (4H, m), 2.80-2.94 (1H, m), 3.01-3.12 (5H, m), 3.94 (2H, m), 4.21 (1H, m), 6.76 (2H, d), 7.09 (2H, d).

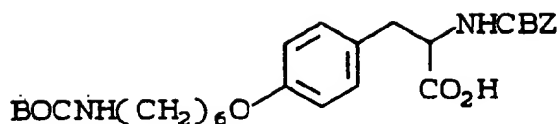
Analysis for $C_{28}H_{35}N_3O_5 \cdot 1.2 H_2O$

Calcd: C, 63.57; H, 7.67; N, 8.56

Found: C, 63.33; H, 7.28; N, 8.55

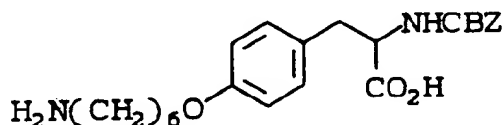
SCHEME 2



EXAMPLE 82-S-(N-(Benzyloxycarbonylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-1)

N-CBZ-L-tyrosine (15.0 g, 0.045 moles) was dissolved in 75 mL DMF and added at 0-10°C to a suspension of sodium hydride (2.16 g, 0.09 moles) in 25 mL DMF. The resulting suspension was stirred at 0-10°C for 1.0 hour and then 6-(t-butyloxycarbonylamino)hexyl bromide (12.6 g, 0.045 moles) in 25 mL DMF was added dropwise at 0-5°C and the clear, dark reaction mixture was stirred at room temperature overnight.

After solvent removal, the residue was taken up in EtOAc and this was made acidic with 10% KHSO₄ solution. The organic phase was separated, washed with brine, dried (Na₂SO₄) and the solvent removed to give an oil. This was purified by column chromatography on silica gel eluting with 98:2:1 CHCl₃/CH₃OH/HOAc to give pure 2-1 as a clear oil. ¹H NMR (300 MHz, CD₃OD) δ 1.45 (15H, m), 1.75 (2H, m), 2.80-3.15 (6H, m), 3.91 (2H, t), 4.38 (1H, m), 4.95 (6H, m), 6.79 (2H, d), 7.10 (2H, d), 7.28 (5H, m).

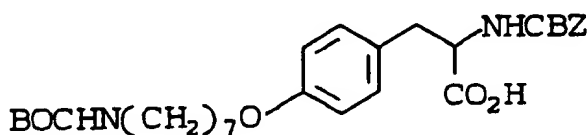
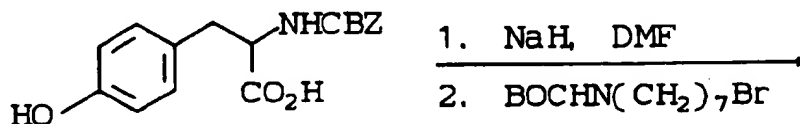
EXAMPLE 92-S-(N-(Benzyloxycarbonylamino)-3-[4-(6-aminohexyloxyphenyl)]propionic acid hydrochloride (2-2)

Compound 2-1 (51.4 mg, 0.1 mmole) was dissolved in 20 mL EtOAc and cooled to -20°C under N₂. HCl gas was bubbled into this solution for 10 minutes as the temperature rose to -5°C. The reaction mixture was stoppered and stirred at 0° to -5°C for 1 hour. The solvent was then removed on the rotary evaporator and the residue was triturated with ether to give 2-2 (14.2 mg) as a white solid. R_f = 0.4 (SiO₂, 9:1:1 EtOH/NH₄OH, H₂O). ¹H NMR (300 MHz, CD₃OD) δ 1.45 (6H, m), 1.73 (4H, m), 2.90 (3H, m), 3.13 (1H, m), 3.95 (2H, m), 4.30 (1H, bs), 6.77 (2H, d), 7.10 (2H, d), 7.32 (4H, m).

Analysis for C₂₃H₃₁N₂O₅Cl·0.5 H₂O

Calcd: C, 60.05; H, 7.01; N, 6.09

Found: C, 60.08; H, 7.06; N, 6.09

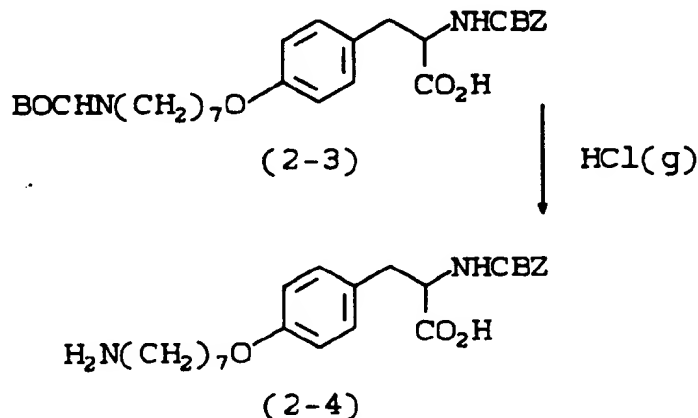
EXAMPLE 102-S-(N-(Benzyloxycarbonylamino)-3-[4-(7-N-t-butyloxycarbonylaminoheptyloxy)phenyl]propionic acid (2-3)

N-CBZ-L-tyrosine (1.27 g, 4.02 mmoles) was alkylated with 7-(N-t-butyloxycarbonylaminoheptyl)bromide

as taught in Example 8 for compound 2-1. Crude product was purified by flash chromatography on silica gel eluting with 95:5:0.5 $\text{CHCl}_3/\text{CH}_3\text{OH}/\text{HOAc}$ to give 1.05 g (50%) of 2-3 as a clear oil.

^1H NMR (300 MHz, CD_3OD) δ 1.40 (20H, m), 1.66 (2H, m), 2.82 (1H, m), 2.97-3.18 (4H, m), 3.91 (2H, m), 4.19 (1H, m), 5.0 (2H, q), 6.77 (2H, d), 7.10 (2H, d), 7.30 (5H, m).

EXAMPLE 11



2-S-(N-(Benzyloxycarbonylamino)-3-[4-(7-aminoheptyloxy)phenyl]propionic acid hydrochloride (2-4)

Compound 2-3 (67.4 mg, 0.127 mmole) was deprotected with HCl gas as described in Example 9 for 2-2 to provide 60.0 mg pure 2-4.

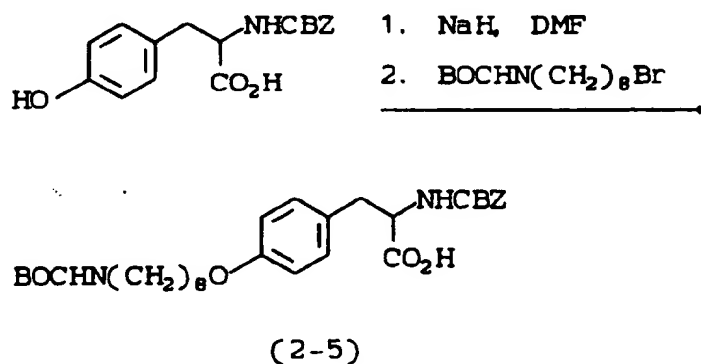
^1H NMR (300 MHz, CD_3OD) δ 1.32 (9H, m), 1.60 (4H, m), 2.77 (3H, m), 3.00 (1H, m), 3.18 (2H, m), 3.72 (2H, m), 4.25 (1H, m), 4.90 (2H, q), 6.70 (2H, d), 7.00 (2H, d), 7.18 (5H, m).

Analysis for $\text{C}_{24}\text{H}_{32}\text{N}_2\text{O}_5 \cdot 0.2\text{EtOH} \cdot 0.75\text{H}_2\text{O}$

Calcd: C, 64.94; H, 7.75; N, 6.21

Found: C, 64.97; H, 7.84; N, 6.22

EXAMPLE 12

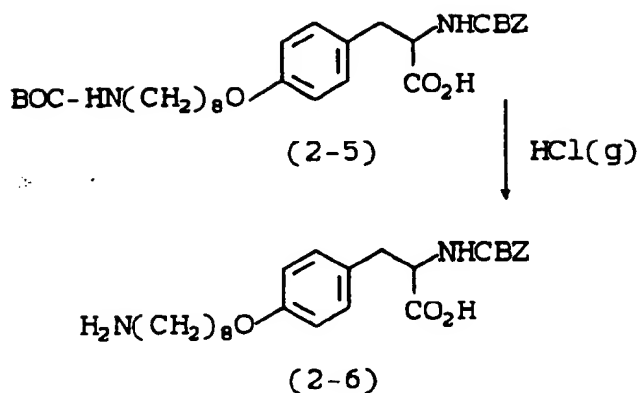


2-S-(N-(Benzyloxycarbonylamino)-3-[4-(8-N-t-butyloxycarbonylamino-octyloxy)phenyl]propionic acid (2-5)

N-CBZ-L-tyrosine- H_2O (1.5 g, 4.29 mmole) was dissolved in $\text{EtOAc}/\text{CH}_2\text{Cl}_2$, dried over MgSO_4 , filtered and evaporated. The residue was dissolved in DMF and treated with NaH (50% dispersion in oil, 0.43 g, 8.96 mmole) for 1 hour. N-BOC-8-amino-1-bromooctan (1.33 g, 4.34 mmole) was added and the reaction was stirred for 16 hours. The DMF was removed *in vacuo*, the residue dissolved in water, acidified to pH 3 and extracted with EtOAc . The EtOAc layers were combined, dried and concentrated. Column chromatography (SiO_2 , 97:3:1 $\text{CHCl}_3/\text{MeOH}/\text{HOAc}$) gave 2-5 (1.35 g) (57% yield).

^1H NMR (300 MHz, CD_3OD) δ 7.3 (m, 5H), 7.1 (d, 2H), 6.78 (d, 2H), 5.0 (2q, 2H), 4.38 dd, 1H), 3.8 (m, 2H), 3.13 (dd, 1H), 3.0 (t, 2H), 2.85 (dd, 1H), 1.75 (m, 2H), 1.4 (s, 9H), 1.35 (m, 3H), 1.3 (bs, 7H).

EXAMPLE 13

2-S-(N-(Benzyloxycarbonylamino)-3-[4-(8-aminooctyloxy)phenyl]propionic acid (2-6)

20 Compound 2-5 (1.35 g, 2.49 mmole) was dissolved in ethyl acetate and treated with HCl gas at -20°C, purged with N₂ and concentrated to give a white solid which was rinsed with ethyl acetate and dried to give 0.965 g of product.

¹H NMR (300 MHz, CD₃OD) δ 7.3 (m, 5H), 7.1 (d, 2H), 6.8 (d, 2H), 5.02, (2q, 2H), 4.35 (dd, 1H), 4.03 (t, 2H), 3.1 (dd, 1H), 2.9 (t, 2H), 2.85 (dd, 1H), 1.75 (m, 2H), 1.65 (m, 2H), 1.5 (m, 2H), 1.4 (bs, 6H).

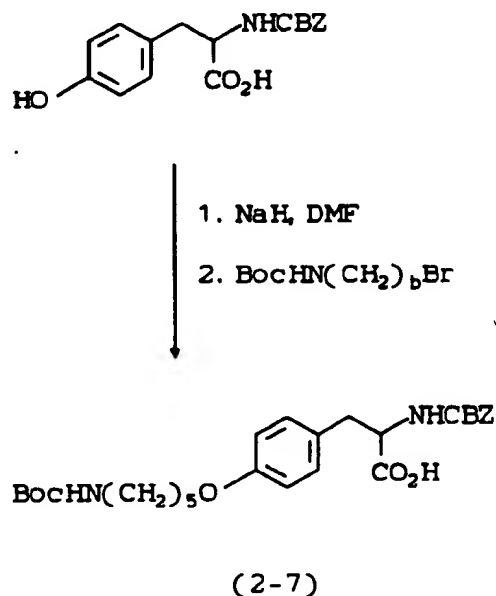
Analysis for C₂₅H₃₄N₂O₅·HCl·0.65 H₂O

25 MW = 490.732

Calcd: C, 61.18; H, 7.46; N, 5.71

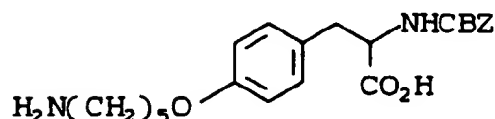
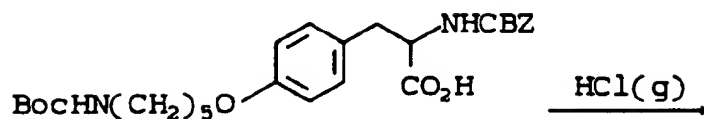
Found: C, 61.18; H, 7.45; N, 5.77

EXAMPLE 14

2-S-(N-(Benzyloxycarbonylamino)-3-[4-(5-N-t-butyloxycarbonylaminopentyloxy)phenyl]propionic acid (2-7)

55 N-CBZ-L-tyrosine (1.06 g, 3.01 mmole) was alkylated with 5-N-t-(butyloxycarbonylaminopentyl) bromide as described for compound 2-1 in Example 8. Crude product was purified by flash chromatography on silica gel eluting with 97:3:0.5 CHCl₃/CH₃OH/HOAc to give pure 2-7.

¹H NMR (300 MHz, CD₃OD) δ 1.42 (9H, s), 1.52 (4H, m), 1.76 (2H, m), 3.05, (3H, m), 3.92 (2H, t), 5.00 (2H, m), 6.79 (2H, d), 7.11 (2H, d), 7.28 (5H, m).

EXAMPLE 15**2-S-(N-(Benzyloxycarbonylamino)-3-[4-(5-amino-pentyloxy)phenyl]propionic acid hydrochloride (2-8)**

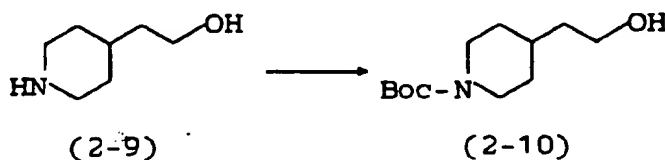
2-7 was treated with HCl gas as taught in Example 9 for compound 2-2, to provide pure 2-8 as a white powder.

¹H NMR (300 MHz, CD₃OD) δ 1.60 (2H, m), 1.77 (4H, m), 2.90 (3H, m), 3.12, (1H, m), 3.96 (2H, t), 4.37 (1H, m), 5.02 (2H, m), 6.81 (2H, d), 7.12 (2H, d), 7.30 (5H, m).

Analysis for C₂₂H₂₉N₂O₅·0.25 H₂O

Calcd: C, 59.85; H, 6.74; N, 6.35

Found: C, 59.85; H, 6.73; N, 6.32

EXAMPLE 16**2-(4-N-t-Butyloxycarbonylpiperidinyl)ethanol (2-10)**

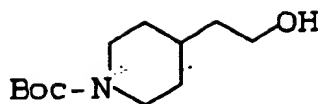
4-piperidine-2-ethanol (Available from American Tokyo Kasei) (130 g, 1.0 mole) was dissolved in 700 mL dioxane, cooled to 0° C and treated with 3 N NaOH (336 mL, 1.0 mole), and di-t-butylcarbonate (221.8 g, 1.0 mole). The ice bath was removed and the reaction stirred overnight. The reaction was concentrated, diluted with water and extracted with ether. The ether layers were combined, washed with brine, dried over MgSO₄, filtered and evaporated to give 225.8 g of product (98%).

R_f = 0.37 in 1:1 EtOAc/Hexanes, ninhydrin stain

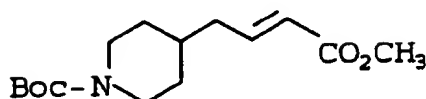
300 MHz ¹H NMR (CDCl₃) δ 4.07 (bs, 2H), 3.7 (bs, 2H), 2.7 (t, J = 12.5 Hz, 2H), 1.8-1.6 (m, 6H), 1.51 (s, 9H), 1.1 (ddd, J = 4.3, 12.5, 12 Hz, 2H).

EXAMPLE 17

1. DMSO, Oxalyl Chloride

2. Carbomethoxytri-
phenylphosphorane

(2-10)

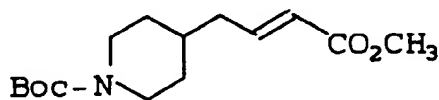


(2-11)

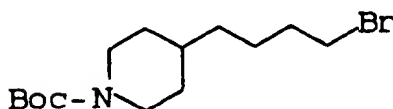
Methyl 4-(4-N-t-Butyloxycarbonylpiperidinyl)-but-2-enoate (2-11)

Oxalyl chloride (55.8 mL, 0.64 mole) was dissolved in 1 L CH_2Cl_2 and cooled to -78°C under N_2 . DMSO (54.2 mL, 0.76 mole) was added dropwise. After gas evolution had ceased, 2-10 (102.5 g, 0.45 mole) dissolved in 200 mL CH_2Cl_2 was added over 20 minutes. After stirring an additional 20 minutes, triethylamine (213 mL, 1.53 mole) was added dropwise and the cold bath removed. After 1 and 1/2 hours TLC showed starting material gone. Carbomethoxytriphenylphosphorane (179 g, 0.536 mole) was added and the reaction stirred overnight. The solution was diluted with 300 mL Et_2O , extracted once with 800 mL H_2O , twice with 300 mL 10% KHSO_4 solution, then once with 300 mL brine. The organic layer was dried over MgSO_4 , filtered and evaporated. Column chromatography (SiO_2 , 5% EtOAc /Hexanes) yielded 78.4 g (62%) of pure 2-11.

300 MHz ^1H NMR (CDCl_3) δ 6.9 (ddd $J = 15.6, 7.6, 7.6$ Hz, 1H), 5.8 (d, $J = 15.6$ Hz, 1H), 4.0 (bs, 2H), 3.7 (s, 3H), 2.6 (t, $J = 12.6$ Hz, 2H), 2.1 (t, $J = 7.4$ Hz, 2H), 1.7-1.4 (m, 3H), 1.4 (s, 9H), 1.1 (m, 2H).

EXAMPLE 181. H_2/Pd on C2. NaOH 3. BH_3 4. $(\text{C}_6\text{H}_5)_3\text{P}$, CBr_4 

(2-11)



(2-12)

4-(4-N-t-Butyloxycarbonylpiperidinyl)butyl bromide (2-12)

Compound 2-11 (36.2 g, 0.128 mole), was dissolved in 500 mL EtOAc. 10% Palladium on carbon (10 g) was added as a slurry in EtOAc and the reaction was placed under H₂ (in a balloon) overnight. The reaction was filtered through Solka-Floc, the cake washed with EtOAc and the ethyl acetate evaporated to give 34.7 g (90%) of 4-(4-N-t-butyloxycarbonylpiperidin-4-yl)butanoate. TLC R_f = 0.69 in 30% EtOAc/Hexanes.

¹H NMR (300 MHz, CDCl₃) δ 4.0 (bs, 2H), 3.6 (s, 3H), 2.60 (t, J = 12.3 Hz, 2H), 2.20 (t, J = 7.4, 2H), 1.6 (m, 4H), 1.40 (s, 9H), 1.40 (m, 1H), 1.20 (m, 2H), 1.0 (m, 2H).

The butanoate ester (45.3 g, 0.159 mole) was dissolved in CH₃OH and treated with 1 N NaOH (500 mL, 0.5 mole) overnight. The solvent was removed in vacuo, water was added and the solution washed with ether, then acidified with 10% KHSO₄ solution. The aqueous layer was washed with ether, the ether layers were combined, washed with brine, dried (MgSO₄), and concentrated to give the corresponding acid as a clear oil (41.85 g, 97% yield).

¹H NMR (300 MHz, CDCl₃) δ 4.0 (bs, 2H), 2.6 (m, 2H), 2.25 (m, 2H), 1.6 (bs, 4H), 1.4 (s, 9H), 1.3-0.9 (9H).

This acid (20.4 g, 0.077 moles) was treated with borane (BH₃/THF, 235 mL, 235 mmole) in THF at 0° for 1 hour. NaOH (1N, 250 mL) was added dropwise and the solution stirred overnight. The reaction was concentrated to remove THF and extracted with ether. The ether extracts were combined, dried over MgSO₄, filtered and evaporated to give the corresponding alcohol as 19.7 g of a colorless oil.

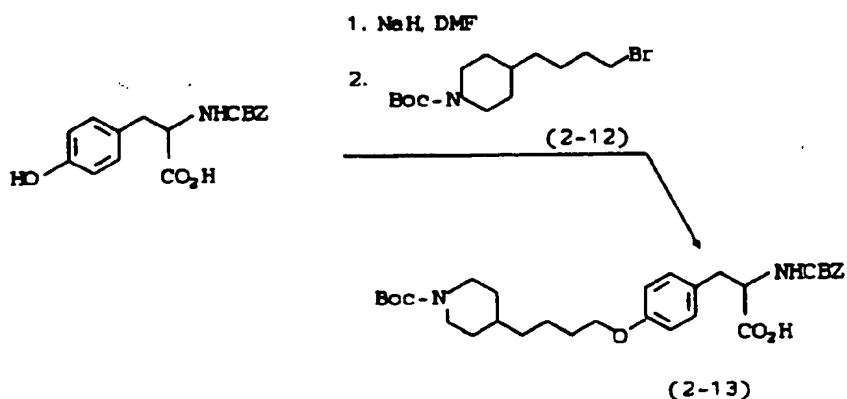
R_f = 0.7 in 2:1 ethyl acetate/hexanes.

¹H NMR (300 MHz, CDCl₃) δ 4.1 (bs, 2H), 3.6 (t, 2H), 2.65 (t, 2H), 2.1 (bs, 1H), 1.65 (bs, 2H), 1.55 (m, 2H), 1.4 (s, 9H), 1.35 (m, 3H), 1.25 (m, 2H), 1.1 (m, 2H).

This alcohol (19.7 g, 76.5 mmole) was dissolved in THF and treated with triphenylphosphine (23.1 g, 88 mmol) and cooled to 0° C. Carbon tetrabromide (29.8 g, 89.9 mmol) was added in one portion, the cold bath was removed and the reaction stirred overnight. Additional triphenyl phosphine (11.71 g) and carbon tetrabromide (14.9 g) was added to drive the reaction to completion. The mixture was filtered and the liquid was diluted with ether and filtered again. After solvent removal the resulting liquid was adsorbed onto SiO₂ and chromatographed with 5% EtOAc/Hexanes to yield 2-12 as a clear colorless oil (20.7 g, 85% yield).

R_f = 0.6 in 1:4 ethyl acetate/hexanes

¹H NMR (300 MHz, CDCl₃) δ 4.1 (bs, 2H), 3.4 (t, 2H), 2.65 (t, 2H), 1.85 (m, 2H), 1.65 (bd, 2H), 1.4 (s, 9H), 1.35 (m, 2H), 1.3 (m, 3H), 1.1 (m, 2H).

EXAMPLE 192-S-(N-(Benzyloxycarbonylamino)-3-[4-(4-N-t-butyloxycarbonylpiperidin-4-yl)butyloxy]phenyl)propionic acid (2-13)

N-CBZ-L-tyrosine was alkylated with 2-12 as taught for compound 2-5 in Example 12 to provide 2-13 in 87% yield.

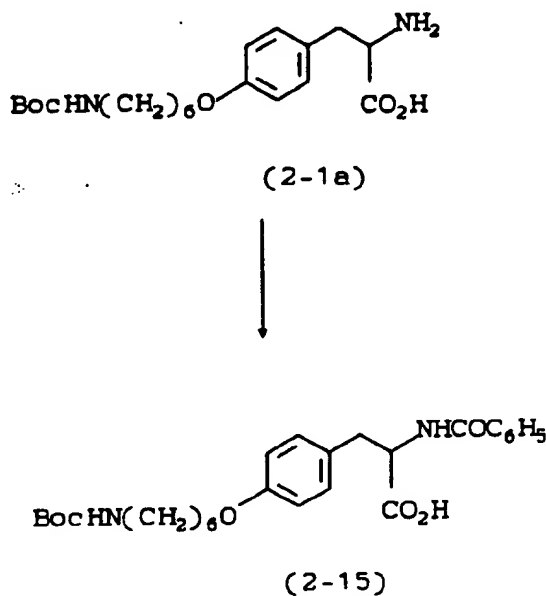
R_f = 0.15 in 97:3:1 CHCl₃/CH₃OH/HOAc, iodine stain.

¹H NMR (300 MHz, CDCl₃) δ 7.2 (d, J = 7.5 Hz, 2H), 7.1 (d, J = 7.5 Hz, 2H), 7.0 (d, J = 7.3 Hz, 2H), 6.8 (d, J = 7.3 Hz, 2H), 5.2 (d, J = 7.9 Hz, 1H), 5.1 (s, 2H), 4.6 (m, 1H), 4.01 (bd, 2H), 3.92 (t, J = 6 Hz, 2H), 6.7 (m, 2H), 2.65 (bt, 7H), 1.75-1.4 (m, 7H), 1.45 (s, 9H), 1.3 (m, 2H), 1.1 (m, 2H).

2-S-Amino-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-1a)

A solution of compound 2-1 (0.52 g, 1.0 mmole) in 20 mL of 4:1 ethanol/HOAc was hydrogenated under balloon pressure for 8 hours. The catalyst was filtered off and the solvent removed on the rotary evaporator to give a residue that was triturated with 30 mL ether to provide 0.16 g of 2-1a.

¹H NMR (300MHz, CD₃OD) δ 1.40 (9H, m), 1.75 (2H, m), 2.90-3.05 (3H, m), 3.10-3.23 (3H, m), 3.70 (1H, m), 3.96 (3H, t), 6.88 (2H, d), 7.20 (2H, d).

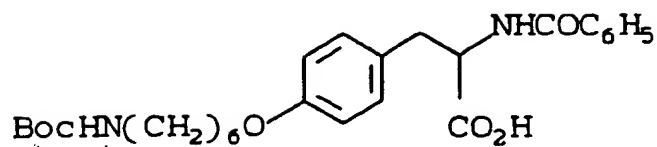
EXAMPLE 222-S-(Phenylcarbonylamino)-3[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl] propionic acid (2-15)

0.152 g (0.4 mmole) of compound 2-1a was added to a solution of 1 N NaOH (0.4 ml) in 10 mL H₂O and this was stirred at 0-5 degrees C for 10 minutes as most of the solid dissolved. To this vigorously stirred suspension was added benzoyl chloride (0.062 g, 0.44 mmole) followed by solid sodium bicarbonate (0.037 g, 0.44 mmol) and the resulting mixture was stirred at 0-5° C for 1 hour.

The reaction mixture was then diluted with 30 mL H₂O and acidified to pH 2-3 with 10% KHSO₄ solution. This was extracted with 3 x 50 mL EtOAc and the combined organic extract was washed with 30 mL of H₂O, 30 mL of brine and dried (Na₂SO₄).

Solvent removal provided a viscous residue that was purified by flash chromatography on silica gel eluting with chloroform(95)-methanol(5) to give 2-15 as a viscous residue.

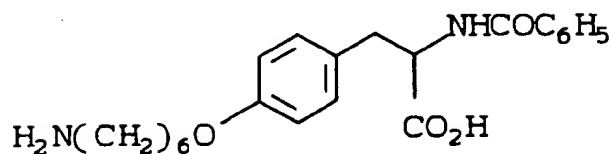
¹H NMR (300MHz, CDCl₃) δ 1.40 (9H, m), 1.75 (2H, bs), 3.20 (m, 4H), 3.92 (2H, m), 5.03 (2H, m), 6.79 (2H,d), 7.10 (2H, d), 7.45 (3H, m), 7.72 (2H, m).

EXAMPLE 23

(2-15)

HCl

EtOAc



(2-16)

2-S-Phenylcarbonylamino-3-[4-(6-aminohexyloxy)phenyl]-propionic acid hydrochloride (2-16)

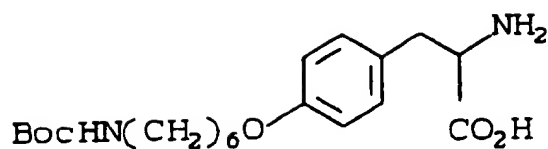
0.28 g (2.0 mmole) of compound 2-15 was dissolved in 20 mL of EtOAc and this was cooled to -15°C and HCl gas was bubbled into the solution for 10 minutes. The resulting mixture was stoppered and stirred at 0°C for 1.5 hours at which point all starting material was consumed. The solvent was then removed on the rotary evaporator to afford a white, foam-like residue. This was stirred with 30 mL ether for 1 hour and the resulting solid was collected by filtration to provide pure 2-16 as a white solid.

^1H NMR (300MHz, CD_3OD), δ 1.50 (3H, m), 1.70 (2H, m), 1.78 (2H, m), 2.90 (2H, t), 3.21 (4H, m), 3.94 (2H, t), 6.80 (2H, d), 7.19 (2H, d), 7.42 (2H, m), 7.50 (1H, m), 7.72 (2H, d).

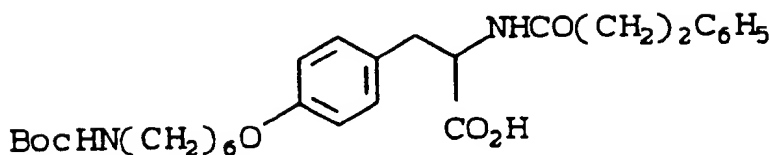
Analysis for $\text{C}_{22}\text{H}_{38}\text{N}_2\text{O}_4\cdot\text{HCl}\cdot 0.75\text{H}_2\text{O}$

Calc.: C = 60.82, H = 6.90, N = 6.45.

Found: C = 60.89, H = 6.67, N = 6.35.

EXAMPLE 24

(2-1 a)

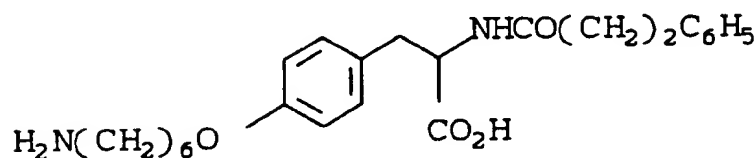
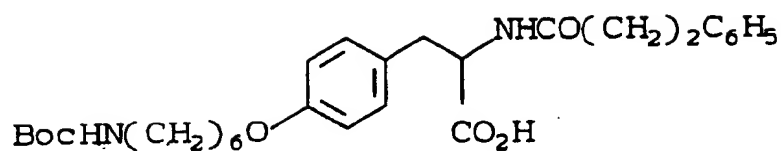


(2-17)

2-S-Phenethylcarbonylamino-3[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propanoic acid (2-17)

To a stirred solution of 1.2 mL 1 N NaOH in 15 mL H₂O cooled to 0-5° C was added 0.457 g (1.2 mmole) of compound 2-1a and the resulting mixture was stirred for 10 minutes during which time most of the solid dissolved. To this vigorously stirred suspension was then added 3-phenylpropanoyl chloride (0.223 g, 1.32 mmole) followed by solid sodium carbonate (0.111 g, 1.32 mmole). The resulting white mixture was stirred vigorously at 0-5° C for 1.5 hours. The reaction mixture was then diluted with 40 mL H₂O and this was acidified to pH 2-3 with a 10% KHSO₄ solution. The resulting aqueous phase was then extracted with 4 x 50 mL portions of EtOAc, and the combined organic phase was washed with 50 mL H₂O, 50 mL brine and dried (Na₂SO₄). Solvent removal gave a viscous solid that was purified by flash chromatography on silica gel, eluting with chloroform (95)-methanol(5) to give 0.30 g of pure 2-17 as a clear viscous gum.

¹H NMR (300 MHz, CDCl₃) δ 1.40 (9H, m), 1.72 (2H, bs), 2.50 (2H, m), 3.02 (6H, m), 3.91 (2H, m), 6.72 (2H, d), 6.88 (2H, m), 7.20 (3H, m), 7.29 (2H, m).

EXAMPLE 252-S-Phenethylcarbonylamino-3-[4-(6-aminohexyloxy)phenyl]propanoic acid hydrochloride (2-18)

30

A solution of compound 2-17 (0.3 g, 3.0 mmole) in 15 mL EtOAc was cooled to -15° C and HCl gas was bubbled in for 10 minutes. The stoppered reaction mixture was then stirred for 2 hours at 0° C at which time all 2-17 was consumed. The solvent was then removed on the rotary evaporator and the resulting foam was triturated with 40 mL ether at room temperature for 1.0 hour to give pure 2-18 as a white solid, 0.22 g.

35

¹H NMR (300 MHz, CD₃OD) δ 1.48 (3H, m), 1.67 (2H, m), 1.80 (2H, m), 2.46 (2H, m), 2.80 (3H, m), 2.90 (2H, m), 3.30 (3H, m), 3.95 (2H, t), 6.79 (2H, d), 7.06 (2H, d), 7.15 (3H, m), 7.22 (2H, m).

Analysis for C₂₄H₃₂N₂O₄·HCl·H₂O

Calc.: C = 61.72, H = 7.55, N = 6.00.

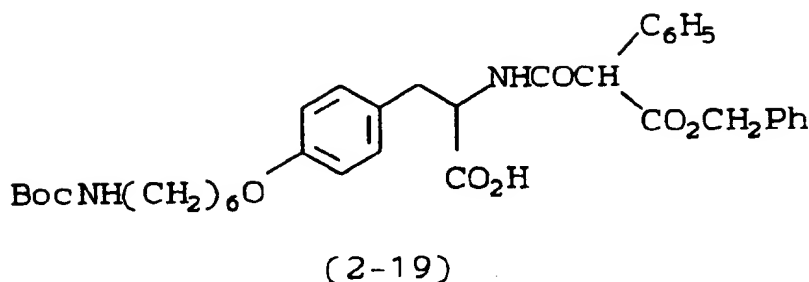
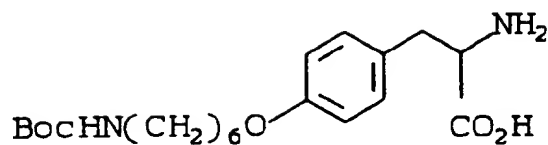
Found: C = 61.97, H = 7.11, N = 5.96.

40

45

50

55

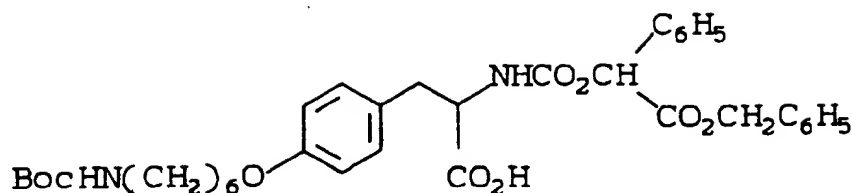
EXAMPLE 26

30

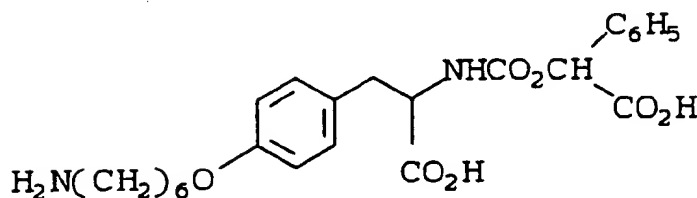
2-S-(2-N-(2-Benzyloxycarbonyl)phenylacetyl)amino-3[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-19)

35 To a cold solution of 1.8 mL of 1 N NaOH in 15 mL H₂O was added 0.685 g (1.8 mmole) of compound 2-1a with stirring to give, after 10 minutes, a clear solution. Then, 2-benzyloxycarbonylphenylacetyl chloride (0.577 g, 2.0 mmole) was added followed by sodium bicarbonate (0.168 g, 2.0 mmole) and the resulting mixture was stirred at 0-5° C for 1.0 hour. The reaction mixture was diluted with water, acidified to pH 2-3 with 10% KHSO₄ solution and extracted with 4 x 500 mL portions of EtOAc. The combined organic extracts were washed with brine, dried (Na₂SO₄) and the solvent was removed to give a viscous amber residue. This was purified by column chromatography on silica gel, eluting with CHCl₃ (98)-methanol (2) to give 0.326 g of pure 2-19 as an oil.

40 ¹H NMR (300 MHz CDCl₃) δ 1.45 (9H, 6s), 1.75 (2H, 6s), 3.07 (4H, m), 3.89 (2H, bs), 4.57 (2H, bs), 5.15 (2H, m), 6.69 (2H, d), 6.88 (2H, d), 7.30 (5H, m).

EXAMPLE 27

(2-19)



(2-20)

2-S-(2-Carboxyphenylacetyl-amino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid hydrochloride (2-20)

Compound 2-19 (0.34 g, 0.55 mmole) was dissolved in 25 mL absolute ethanol and after adding 100 mg 10% Pd/C the suspension was hydrogenated under balloon pressure. Then, the catalyst was filtered off and the solvent removed on the rotary evaporator to give 0.25 g of 2-S-(2-Carboxyphenylacetyl-amino)-3-[4-(6-t-butyloxycarbonylamino-hexyloxy)phenyl]propionic acid.

^1H NMR (300MHz, CD_3OD) δ 1.47 (12H, m), 1.78 (2H, m), 3.06 (3H, m), 3.32 (4H, m), 3.92 (2H, m), 4.60 (2H, m), 6.72 (2H, d), 6.96, (2H, d), 7.30 (5H, m).

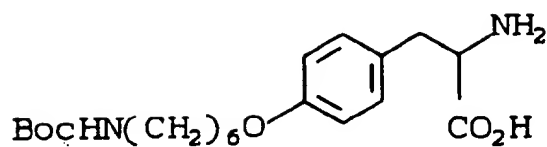
This acid was dissolved in 25 mL EtOAc and treated with HCl gas as described for compound 2-2 in Example 9. Solvent removal provided a residue that was purified by flash chromatography on silica gel eluting with 9:1:1 ethanol/ H_2O / NH_4OH to give pure 2-20.

^1H NMR (300 MHz, D_2O) δ 1.55 (H, m), 1.90 (2H, m), 2.83-3.09 (4H, m), 3.28 (1H, m), 4.15 (2H, m), 6.887.45 (9H, m).

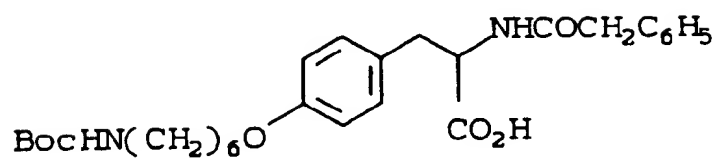
Analysis for $\text{C}_{24}\text{H}_{30}\text{N}_2\text{O}_6 \cdot 1.5 \text{ H}_2\text{O} \cdot 0.25 \text{ NH}_3$

Calc.: C = 60.84, H = 7.18, N = 6.65.

Found: C = 60.48, H = 6.81, N = 6.99.

EXAMPLE 28

(2-1a)

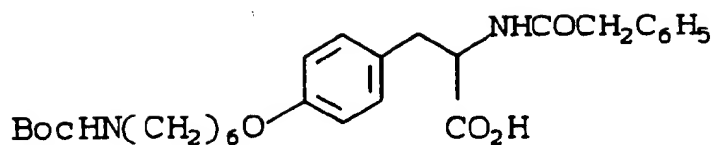


(2-21)

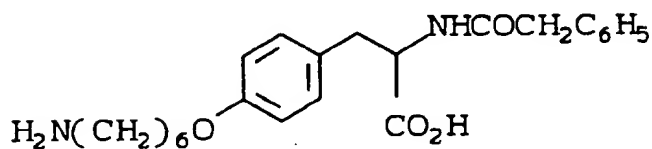
2-S-(Phenylacetyl-amino)-3-[4-(6-N-t-butyloxycarbonylamino-hexyloxy)phenyl]propionic acid (2-21)

Compound 2-1a (0.685 g, 1.8 mmole) was acylated with phenylacetyl chloride as described for compound 2-19 in Example 26. The crude product was purified by flash chromatography on silica gel eluting with 95:5:0.5 CHCl₃/CH₃OH/HOAc to give pure 2-21 as a viscous oil. (0.35 g).

¹H NMR (300 MHz, CD₃OD) δ 1.45 (12H, m), 1.78 (2H, m), 2.88 (1H, m), 3.10 (3H, m), 3.30 (1H, m), 3.48 (2H, m), 3.92 (2H, m), 4.61 (1H, m), 6.74 (2H, d), 7.02 (2H, d), 7.12 (2H, m), 7.22 (3H, m).

EXAMPLE 29

(2-21)



(2-22)

2-S-(Phenylacetylamino)-3-[4-(6-aminohexyloxy)phenyl]-propionic acid (2-22)

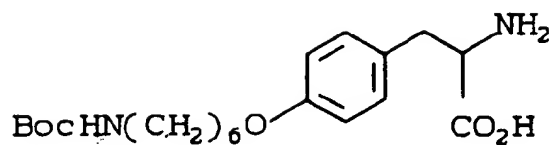
Compound 2-21 (0.35 g) was dissolved in 25 mL EtOAc and this solution was treated with HCl gas as described for compound 2-16 in Example 23 to give 0.26 g pure 2-22 as a white solid.

¹H NMR (300 MHz, CD₃OD) δ 1.50 (6H, m), 1.65 (2H, m), 2.20 (2H, m), 2.88 (3H, m), 3.12 (1H, m), 3.30 (2H, m), 3.47 (2H, m), 3.94 (2H, m), 4.61 (1H, m), 6.75 (2H, d), 7.02 (2H, d), 7.13 (2H, d), 7.30 (3H, m).

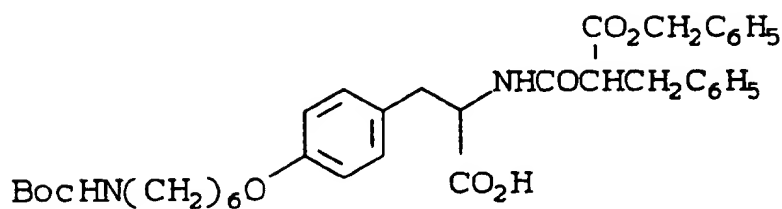
Analysis for C₂₃H₃₀N₂O₄·HCl·H₂O

Calc.: C = 60.98, H = 7.34, N = 6.19.

Found: C = 61.29, H = 6.92, N = 6.12.

EXAMPLE 30

(2-1a)



(2-23)

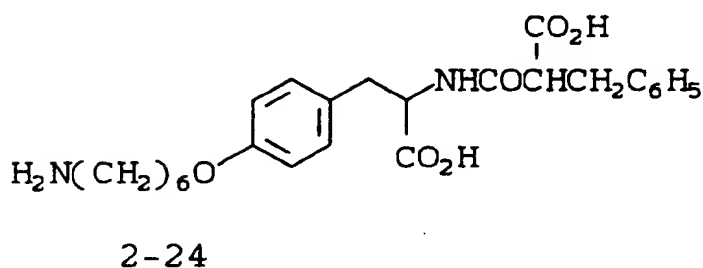
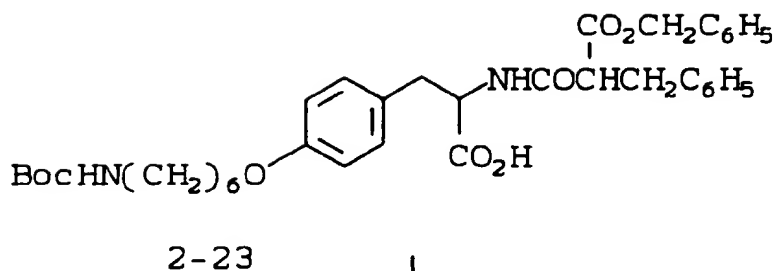
25

30

2-S-[(2-N-Benzyloxycarbonyl-3-phenylpropionylamino]-3[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-23)

Compound 2-1a (0.685 g, 1.8 mmole) was acylated with 2-N-benzyloxycarbonyl-3-phenylpropionylchloride as described for compound 2-19 in Example 26. The crude product was purified by flash chromatography on silica gel eluting with 98:2:1 CHCl₃/CH₃OH/HOAc to give pure 2-23 as a viscous oil.

35 ¹H NMR (300 MHz, CD₃OD) δ 1.40 (16 H, m), 1.61 (2H, m), 3.03 (8H, m), 3.30 (6H, m), 3.71 (1H, m), 3.86 (2H,m), 4.60 (1H, m), 5.02 (2H, m), 6.70 (2H, d), 6.86, (1H, d), 7.02 (1H, 3), 7.22 (5H, m).

EXAMPLE 31**2-S-(2-Carboxy-3-phenylpropionylamino)-3-[4(6-aminohexyloxy)phenyl]propionic acid (2-24)**

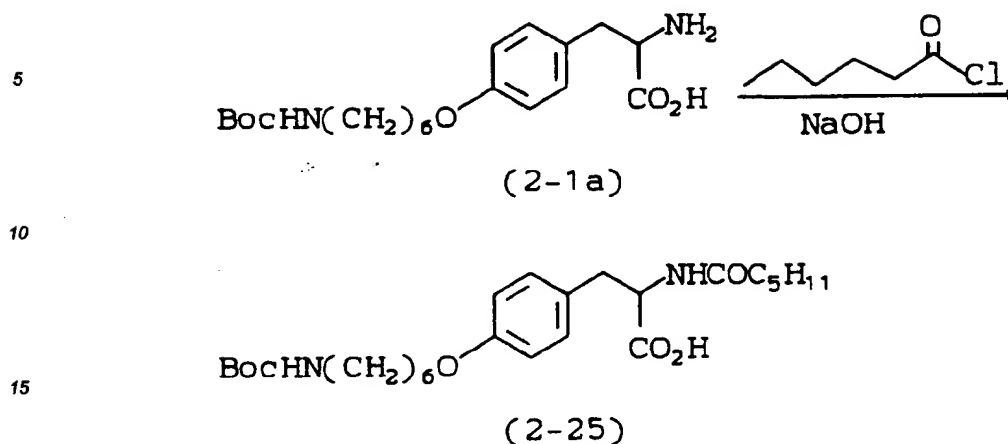
30 Compound 2-23 (0.49 g, 0.76 mmole) was dissolved in 25 mL ethanol and after the addition of 100 mg 10% Pd/C was hydrogenated at balloon pressure overnight. Solvent removal provided 2-S-(2-carboxy-3-phenylpropionylamino)-3-[4-(6-N-t-butyloxycarbonylamino)hexyloxy]phenyl]propionic acid as a viscous residue (0.35 g).

¹H NMR (300 MHz, CD₃OD) δ 1.42 (10H, m), 1.75 (2H, m), 2.80-3.15 (5H, m), 3.30 (1H, m), 3.90 (2H, m), 4.58 (2H, m), 6.68-6.85 (4H, m), 7.06-7.27 (5H, m).

35 This acid (0.32 g) was treated with HCl gas as described for compound 2-12 to give after solvent removal a crude product that was purified by flash chromatography on silica gel eluting with 90:5:5 CHCl₃/CH₃OH/HOAc to provide the diastereomeric products 2-24a and 2-24b.

2-24a had ¹H NMR (300 MHz, D₂O) δ 1.58 (4H, m), 1.83 (4H, m), 2.95 (2H, m), 3.08 (3H, m), 3.20 (1H, m), 3.51 (1H, m), 4.18 (2H, m), 4.53 (1H, m), 4.95 (2H, g), 6.92 (4H, m), 7.43 (5H, m).

40 2-24b had ¹H NMR (400 MHz, D₂O) δ 1.40 (4H, m), 1.62 (2H, m), 1.73 (2H, m), 2.90 (6H, m), 3.31 (1H, m), 4.17 (2H, m), 4.32 (1H, m), 6.93 (2H, d), 7.07 (2H, d), 7.15 (2H, d), 7.26 (3H, m).

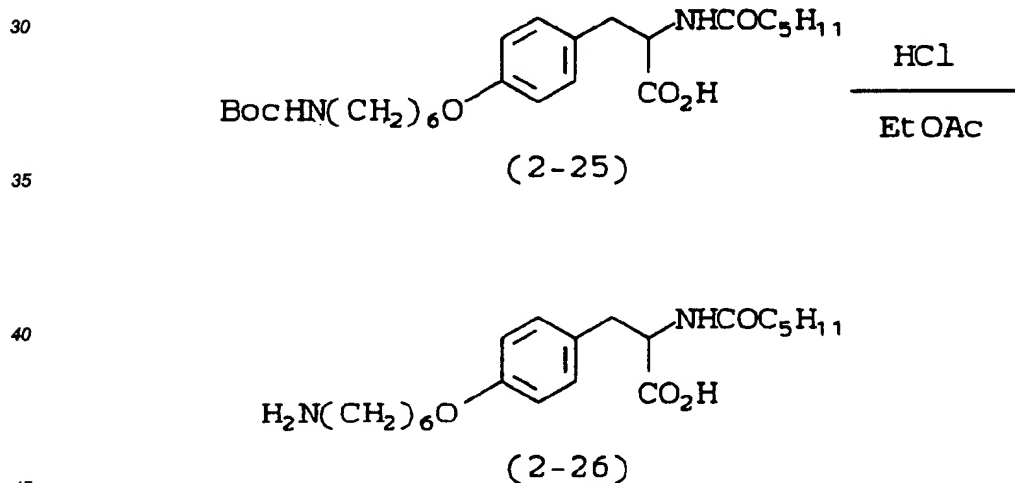
EXAMPLE 31(a)2-S-(Hexanoylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-25)

20

2-1a (0.685 g, 1.8 mmole) was treated with hexanoyl chloride (0.38 g, 2.0 mmole) as described for 2-15 to provide crude 2-25. This was purified by flash chromatography on silica gel eluting with 95:5:1 CHCl₃/CH₃OH/HOAc to give pure 2-25 as an oil (0.35 g, 41%).

25

¹H NMR (300 MHz, CDCl₃) δ 0.89 (3H, t), 1.20-1.65 (21H, m), 1.75 (2H, m), 2.19 (2H, t), 3.11 (4H, m), 3.92 (2H, m), 4.83 (1H, m), 6.80 (2H, d), 7.05 (2H, d).

EXAMPLE 31(b)2-S-(Hexanoylamino)-3-[4-(6-aminohexyloxy)phenyl]-propionic acid hydrochloride (2-26)

2-25 (0.35 g, 0.75 mmole) was dissolved in 30 mL EtOAc and treated with HCl as described for compound 2-2 to give a foam-like solid that was triturated with 50 mL of ether for 1 hour at room temperature. This gave pure 2-26 as a white solid. (0.186 g).

50

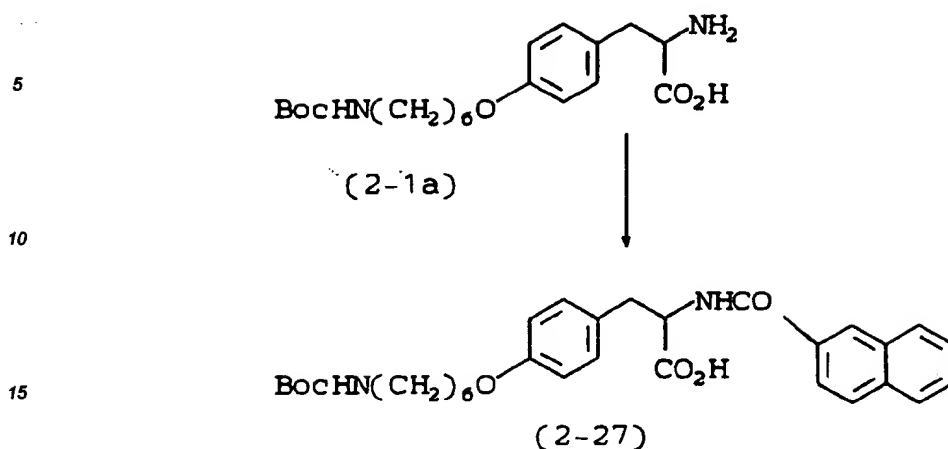
¹H NMR (300 MHz, CD₃OD) δ 0.85 (3H, t), 1.20 (4H, m), 1.48 (6H, m), 1.68 (2H, m), 1.77 (2H, m), 2.14 (2H, m), 4.61 (1H, m), 6.80 (2H, d), 7.13 (2H, m).

Analysis for C₂₁H₃₄N₂O₄·HCl·0.5 H₂O

55

Calc: C=59.49, H=8.56, N=6.61

Found: C=59.32, H=8.48, N=6.55

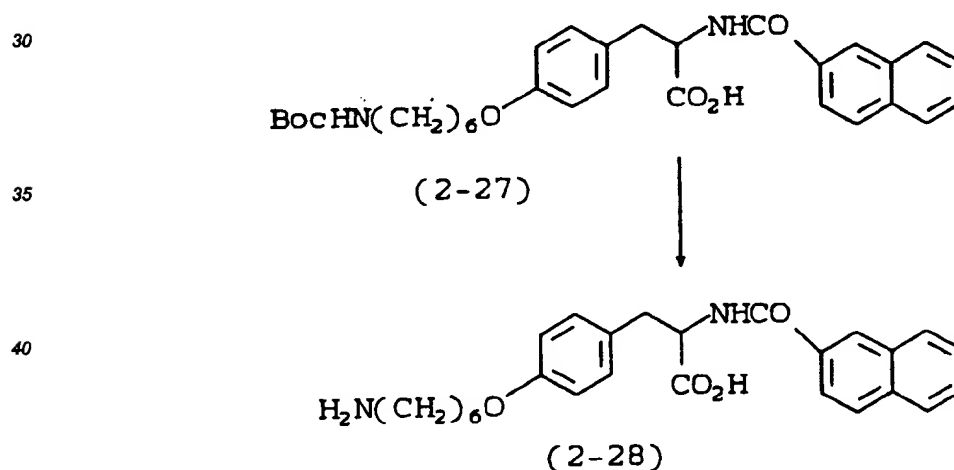
EXAMPLE 31(c)2-S-(2-Napthanoylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-27)

20

2-1a (0.685 g, 1.8 mmole) was treated with 2-napthanoyl chloride (0.409 g, 2.0 mmole) as described for 2-15 to provide crude 2-27. This was purified by flash chromatography on silica gel eluting with 95:4:1 CHCl₃/CH₃OH/HOAc to give pure 2-27 as a white solid (0.14 g).

25

¹H NMR (300 MHz, CD₃OD) δ 1.45 (16H, m), 1.70 (2H, m), 2.88 (1H, m), 3.08 (3H, m), 3.57-3.80 (4H, m), 4.62 (1H, m), 6.54 (2H, d), 6.92 (2H, d), 7.25 (1H, d), 7.42 (2H, m), 7.61 (1H, bs), 7.77 (3H, m).

EXAMPLE 31(d)2-S-(Naphthanoylamino)-3-[4-(6-aminohexyloxy)phenyl]-propionic acid (2-28)

2-27 (0.14 g, 0.31 mmole) was dissolved in 25 mL EtOAc and treated with HCl gas as described for 2-2. Crude product was purified by flash chromatography on silica gel eluting with 10:1:1 C₂H₅OH/H₂O/NH₄OH to give pure 2-28 (55 mg) as a white solid.

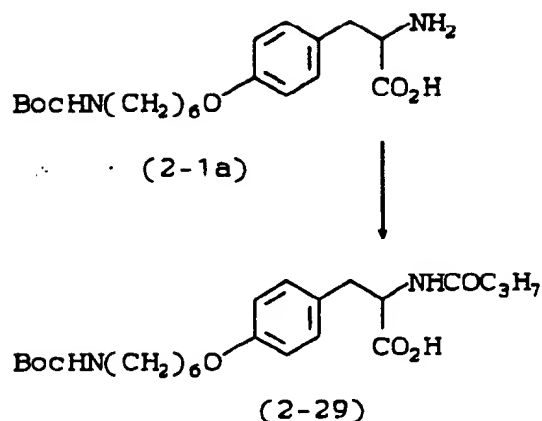
50

¹H NMR (300 MHz, CD₃OD), δ 1.42 (5H, m), 1.71 (2H, m), 2.63 (2H, m), 2.86 (1H, m), 3.07 (2H, m), 3.30 (3H, m), 3.55-3.75 (4H, m), 4.47 (1H, m), 6.43 (2H, d), 6.82 (2H, d), 7.30 (1H, dd), 7.45 (2H, m), 7.64 (1H, bs), 7.80 (3H, m).

Analysis for C₂₇H₃₂N₂O₄·0.5 H₂O

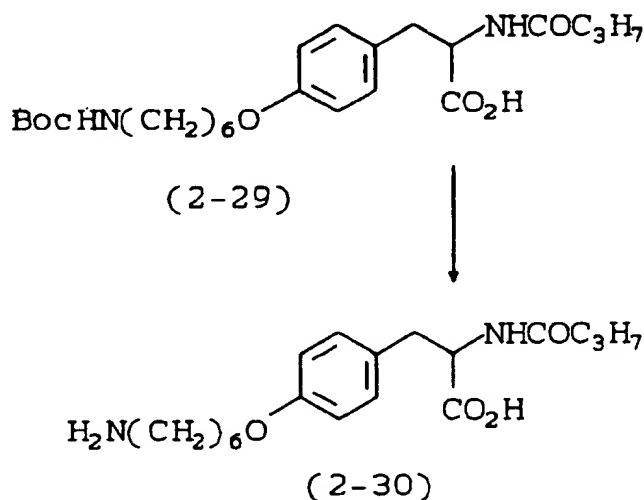
55 Calc.: C=70.87, H=7.27, N=6.12

Found: C=70.93, H=7.04, N=6.11

EXAMPLE 31(e)2-S-(2-Butanoylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-29)

20 2-1a (0.685 g, 1.8 mmole) was acylated with butanoyl chloride as described for 2-15 to give crude 2-29. This was purified by flash chromatography eluting with 95:4:1 CHCl₃/CH₃OH/HOAc to provide pure 2-29 as an oil.

¹H NMR (300 MHz, CD₃OD) δ 0.73 (3H, t), 1.32-1.60 (16H, m), 1.73 (2H, m), 2.12 (2H, m), 2.87 (1H, m), 3.03 (2H, t), 3.12 (1H, m), 3.92 (2H, t), 4.61 (1H, m), 6.80 (2H, d), 7.12 (2H, d).

EXAMPLE 31(f)2-S-(Butanoylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid (2-30)

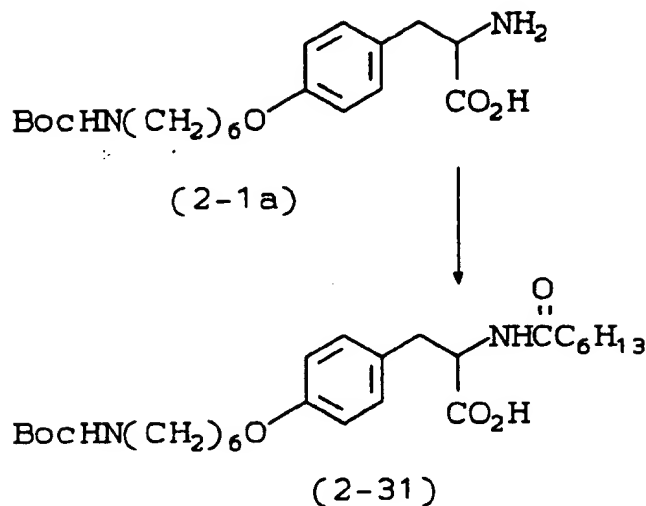
2-29 (0.05 g, 1.0 mmole) was dissolved in 25 mL ethyl acetate and treated with HCl gas as described for 2-2. Crude reaction product was triturated with 25 mL ether to give pure 2-30 as a white solid.

50 ¹H NMR (300 MHz, CD₃OD) δ 0.72 (3H, t), 1.45-1.60 (6H, m), 1.70 (2H, m), 1.79 (2H, m), 2.12 (2H, m), 2.80-2.95 (3H, m), 3.14 (1H, dd), 3.30 (1H, m), 3.95 (2H, t), 4.40 (1H, m), 6.80 (2H, d), 7.13 (2H, d).

Analysis for C₁₉H₃₀N₂O₄·HCl·H₂O

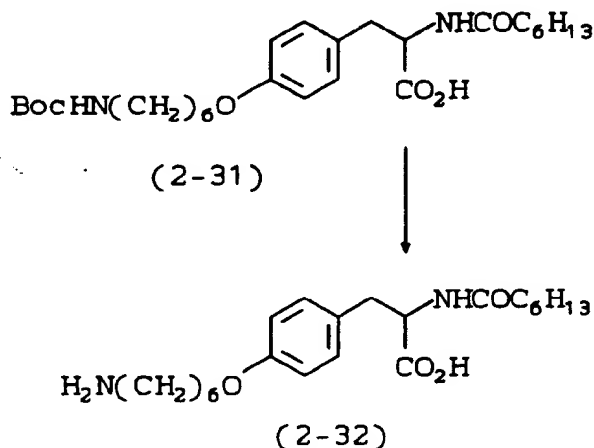
Calc.: C = 56.35, H = 8.21, N = 6.92

Found: C = 56.70, H = 8.12, N = 6.91.

EXAMPLE 31(g)2-S-(Heptanoylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-31)

2-1a (0.685 g, 1.8 mmole) was acylated with heptanoyl chloride as described for 2-15. Crude product was purified by flash chromatography on silica gel eluting with 96:3:1 CHCl₃/CH₃OH/HOAc to give pure 2-31 (0.07 g) as an oil.

¹H NMR (300 MHz, CD₃OD) δ 0.78 (3H, t), 1.22 (6H, m), 1.32-1.55 (16H, m), 1.73 (2H, m), 2.13 (2H, m), 2.85 (1H, m), 3.02 (2H, t), 3.15 (1H, m), 4.91 (2H, t), 4.61 (1H, m), 6.81 (2H, d), 7.12 (2H, d).

EXAMPLE 31(h)2-S-(Heptanoylamino)-3-[4-(6-aminohexyloxy)phenyl]-propionic acid hydrochloride (2-32)

2-31 (0.070 g) was dissolved in 30 mL EtOAc and treated with HCl gas as described for 2-2. Crude reaction product was triturated with 30 mL ether to provide pure 2-32 (52 mg) as a white solid.

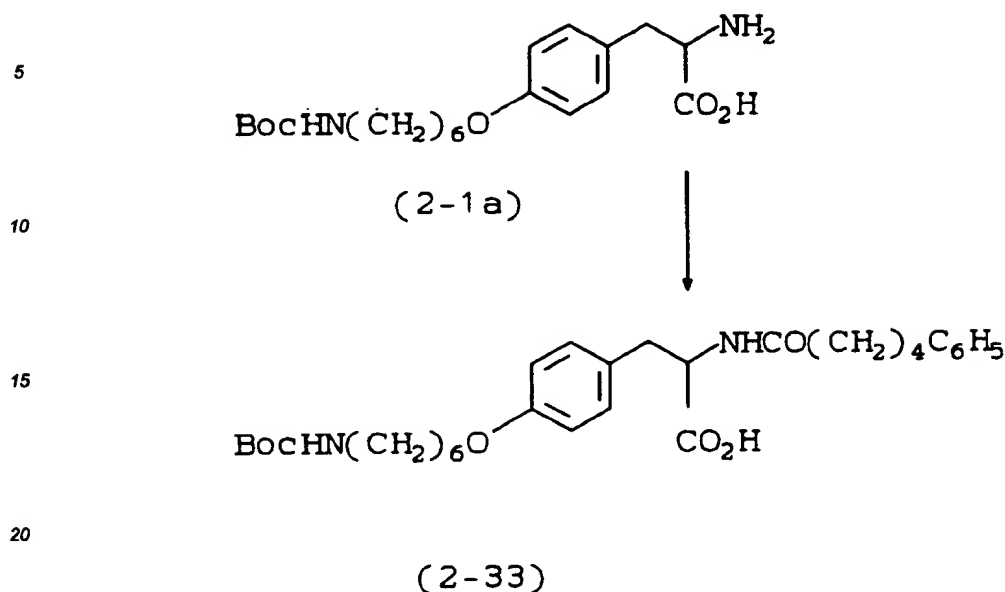
¹H NMR (300 MHz, CD₃OD) δ 0.88 (3H, t), 1.22 (6H, m), 1.47 (6H, m), 1.68 (2H, m), 1.78 (2H, m), 2.13 (2H, t), 2.80-2.95 (3H, m), 3.14 (1H, m), 3.30 (1H, m), 3.94 (2H, m), 4.61 (1H, m), 6.80 (2H, d), 7.13 (2H, d).

Analysis for C₂₂H₃₈N₂O₄·HCl·0.75 H₂O

Calc.: C = 59.71, H = 8.77, N = 6.33

Found: C = 59.76, H = 8.40, N = 6.25.

EXAMPLE 31(i)

2-(S)-(5-Phenylpentanoylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (2-33)

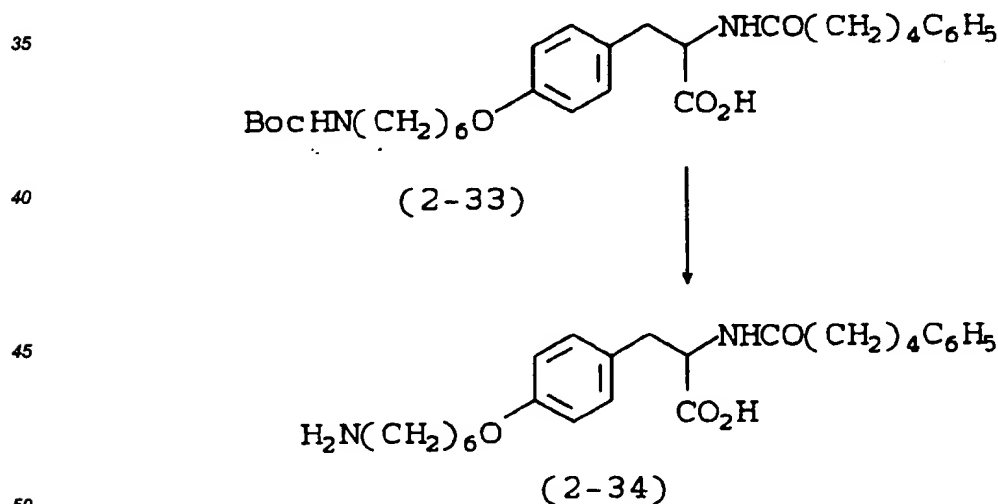
25

2-1a (0.685 g, 1.8 mmole) was acylated with 5-phenylpentanoyl chloride as described for 2-15. Crude product was purified by flash chromatography on silica gel eluting with 96:3:1 CHCl₃/CH₃OH/HOAc to give pure 2-33 (0.49 g) as a clear oil.

¹H NMR (300 MHz, CD₃OD) δ 1.32-1.60 (1H, m), 1.73 (2H, m), 2.18 (2H, m), 2.53 (2H, m), 2.80-2.90 (1H, m), 3.02 (2H, t), 3.04 (1H, m), 4.62 (1H, m), 6.78 (2H, d), 7.08-7.28 (7H, m).

30

EXAMPLE 31(j)

2-S-(5-Phenylpentanoylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid hydrochloride (2-34)

55

2-33 (0.49 g) was dissolved in 30 mL ethyl acetate and treated with HCl gas as described for 2-2. Crude product was triturated with 50 mL ether to give pure 2-34 (0.32 g) as a white solid.

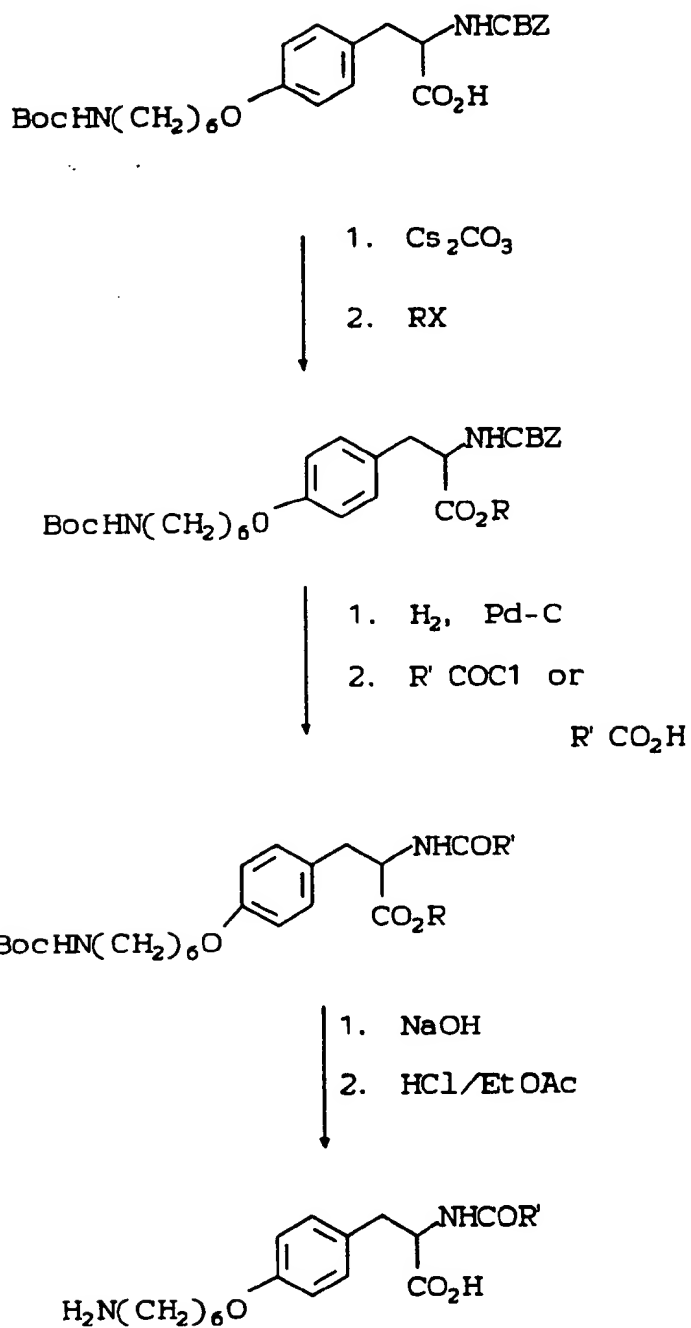
¹H NMR (300 MHz, CD₃OD) δ 1.40-1.58 (8H, m), 1.62-1.70 (2H, m), 1.80 (2H, m), 2.19 (2H, m), 2.55 (2H, m), 2.80-2.95 (3H, m), 3.15 (1H, m), 3.30 (1H, m), 3.90 (2H, t), 4.62 (1H, m), 6.88 (2H, d), 7.08-7.27 (7H, m).

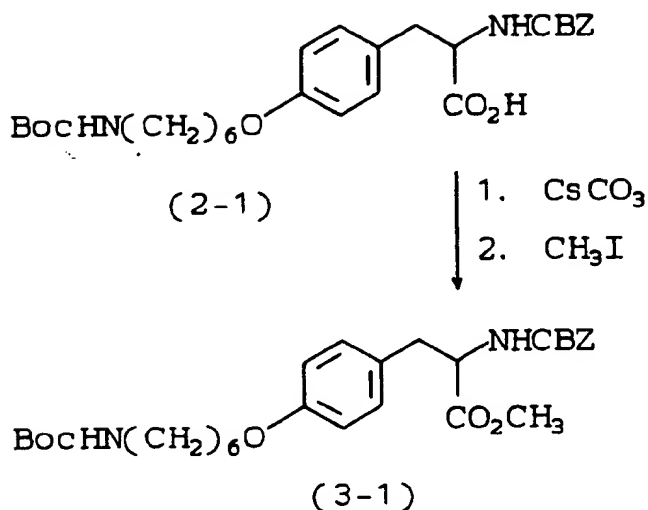
Analysis for C₂₈H₃₈N₂O₄·HCl·H₂O

Calc.: C = 64.24, H = 7.88, N = 5.76

Found: C = 64.53, H = 7.84, N = 5.71.

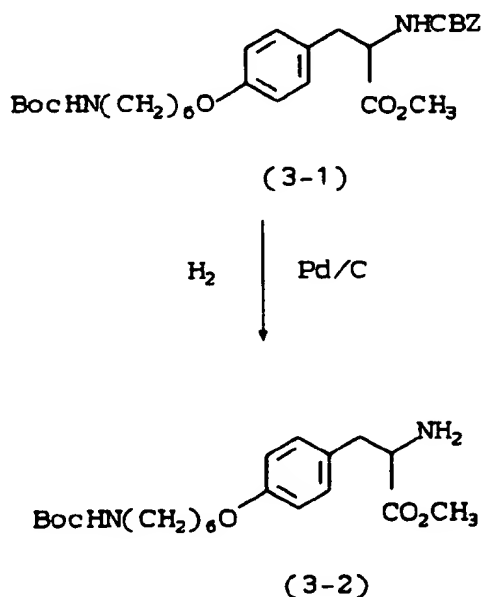
SCHEME 3



EXAMPLE 32**Methyl 2-S-(N-Benzyloxycarbonylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyl)oxyphenyl]propionate (3-1)**

Compound 2-1 (10.0 g, 19.43 mmole) in 75 mL DMF was treated with cesium carbonate (3.16 g, 9.72 mmole) with stirring at room temperature for 2.0 hours. Then, methyl iodide (2.76 g, 19.43 mmole) was added dropwise and the reaction mixture was stirred overnight at ambient temperature. The solvent was removed at high vacuum (30 degrees C) and the residue was taken up in 300 mL EtOAc and washed with 2x40 mL portions of saturated NaHCO₃ solution, brine, and dried (Na₂SO₄). Solvent removal provided 3-1 (8.5 g, 83%) as a clear oil.

¹H NMR (300 MHz, CDCl₃) δ 1.25-1.53 (16H, m), 1.76 (2H, m), 2.96-3.17 (4H, m), 3.71 (3H, s), 3.90 (2H, t), 4.61 (1H, m), 5.10 (2H, m), 5.19 (1H, m), 6.88 (2H, d), 6.98 (sH, d), 7.32 (5H, m).

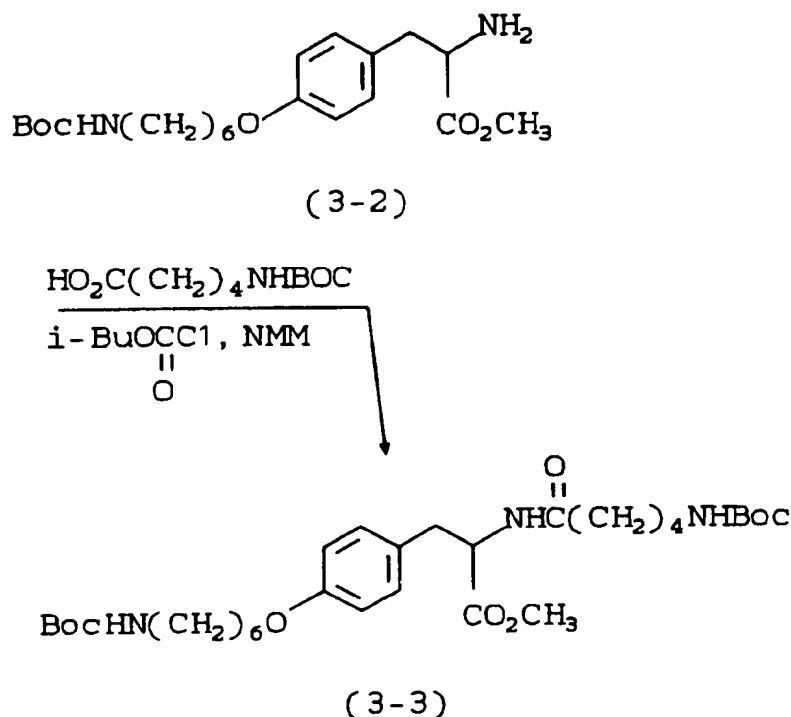
EXAMPLE 33**Methyl 2-S-Amino-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionate (3-2)**

Compound 3-1 (8.0 g, 15.1 mmole) was dissolved in 150 mL absolute ethanol and 1.0 g 10% Pd/C was added. This suspension was hydrogenated in a Parr apparatus (50 psi) for 3.5 hours. The catalyst was then

filtered off and the solvent removed on the rotary evaporator to give pure 3-2 (5.56 g) as a clear oil. $R_f = 0.4$ on SiO_2 with 95:5 $\text{CHCl}_3/\text{CH}_3\text{OH}$

^1H NMR (300 MHz, CDCl_3) δ 1.30-1.55 (16 H, m), 1.70 (2H, m), 2.80 (1H, m), 3.00-3.17 (3H, m), 3.71 (3H, s), 3.93 (2H, t), 6.82 (2H, d), 7.09 (2H, d).

EXAMPLE 34

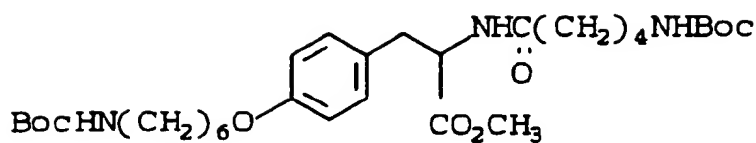


Methyl 2-S-[(5-N-t-Butyloxycarbonylamino)pentanoylamino]-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionate (3-3)

To a solution of 5-(N-t-butyloxycarbonylamino)pentanoic acid (0.293 g, 1.35 mmole) and N-methyl-morpholine (0.187 g, 1.35 mmole) in 10 mL EtOAc at 0-5° C was added i-butylchloroformate (0.184 g, 1.35 mmole) via syringe and the resulting white suspension was stirred for 0.5 hours. Then, 3-2 (0.5 g, 1.27 mmole) dissolved in 10 mL EtOAc was added dropwise and the reaction mixture was stirred at 0° C for 2.0 hours. The reaction mixture was then diluted with 25 mL water/ 40 mL EtOAc and the organic phase was separated, washed with water, 10% KHSO_4 , water, saturated NaHCO_3 , brine and dried (Na_2SO_4). Solvent removal gave an oil that was purified by flash chromatography on silica gel eluting with 2% $\text{CH}_3\text{OH}/\text{CHCl}_3$ ($R_f = 0.35$) to give pure 3-3 (0.68 g, 90%) as a clear oil.

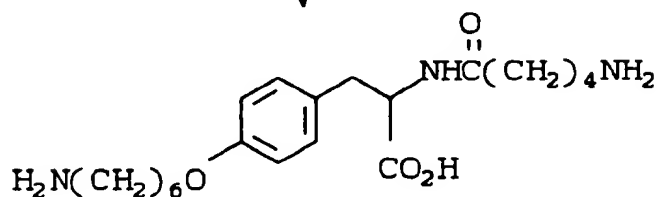
^1H NMR (300 MHz, CDCl_3) δ 1.35-1.55 (26H, m) 1.62 (2H, m), 1.68 (2H, m), 2.20 (2H, t), 3.0-3.16 (6H, m), 3.33 (3H, s), 3.92 (2H, t), 4.83 (1H, m), 6.80 (2H, d), 6.99 (2H, m).

EXAMPLE 35



(3-3)

1. LiOH
2. HCl, EtOAc



(3-4)

2-S-(5-Aminopentanoyl)amino-3-[4-(6-aminohexyloxy)-phenyl]propionic acid dihydrochloride (3-4)

3-3 (0.68 g, 1.14 mmole) was dissolved in 30 mL THF(1)/H₂O(1)/CH₃OH(1), LiOH (0.137 g, 5.73 mmole) was added and the reaction mixture stirred at room temperature overnight. The solvent was then removed and the residue was taken up in 75 mL H₂O and acidified to pH 2-3 with 10% KHSO₄ solution. This was extracted with EtOAc and the combined organic extracts were washed with brine and dried (Na₂SO₄). Solvent removal gave 2-S-(5-t-butyloxycarbonylamino-pentyl)amino-3-[4-(6-t-butyloxycarbonylamino-hexyl)oxyphenyl]-propionic acid (0.65 g).

¹H NMR (300 MHz, CDCl₃) δ 1.40-1.55 (22H, m), 1.60 (2H, m), 1.73 (2H, m), 2.20 (2H, m), 3.10 (4H, m), 3.90 (2H, m), 4.60 (1H, m), 4.72 (1H, m), 4.83 (1H, m), 6.78 (2H, d), 7.05 (2H, d).

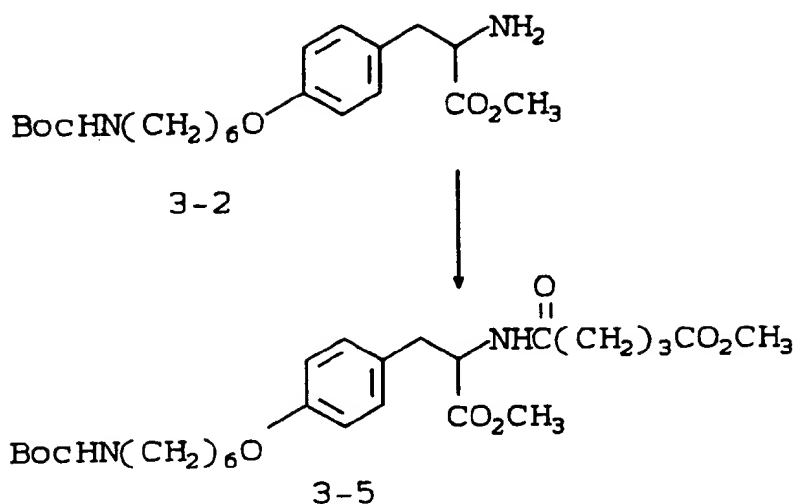
This acid was dissolved in EtOAc and was treated with HCl gas as described for 2-2. The crude hygroscopic white solid was triturated with a solution of 10 mL EtOAc/50 mL Et₂O to give pure 3-4 as a white solid.

¹H NMR (300 MHz, CD₃OD) δ 1.42-1.85 (14H, m), 2.23 (2H, m), 2.90 (6H, m), 3.14 (1H, dd), 3.30 (1H, m), 3.97 (2H, t), 4.60 (1H, m), 6.82 (2H, d), 7.13 (2H, d).

Analysis for C₂₀H₃₃N₃O₄·2HCl·3H₂O

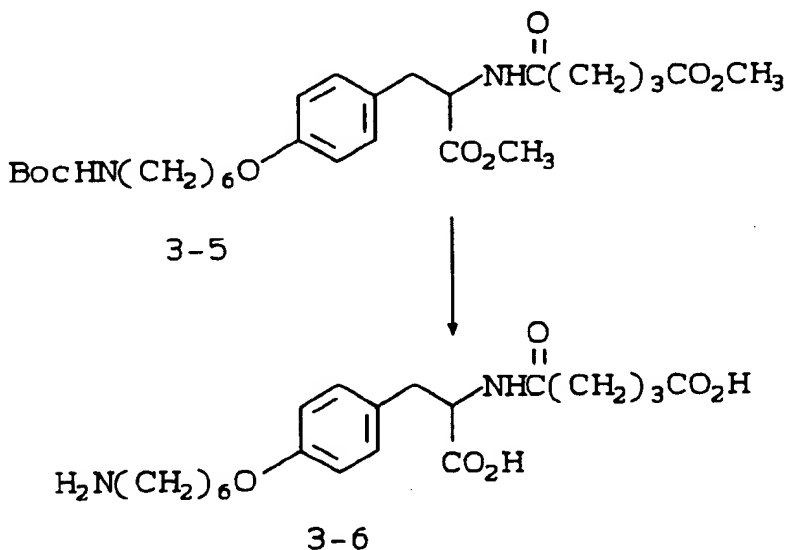
Calc.: C = 47.43, H = 8.16, N = 8.30

Found: C = 47.87, H = 7.49, N = 7.90

EXAMPLE 36**Methyl 2-S-(4-Carbomethoxybutanoyl)amino-3-[4-(N-t-butyloxycarbonylaminohexyloxy)phenyl]propionate (3-5)**

25 To a solution of 3-2 (0.5 g, 1.27 mmole), 4-carbomethoxybutanoic acid (0.213 g, 1.5 mmole) and 1 drop of triethylamine in 20 mL CH₃CN was added BOP reagent (0.66 g, 1.5 mmole) and the resulting clear solution was stirred overnight at room temperature. The solvent was removed on the rotary evaporator and the residue was taken up in EtOAc and this was washed with H₂O, 10% KHSO₄, H₂O, saturated NaHCO₃, brine and dried (Na₂SO₄). Solvent removal provided a residue that was purified by flash chromatography on silica gel eluting with 1% CH₃OH/CHCl₃ to give pure 3-5 (110 mg) as a clear oil.

30 ¹H NMR (300 MHz, CDCl₃), δ 1.35-1.55 (14H, m), 1.75 (3H, m), 1.94 (2H, m), 2.26 (2H, t), 2.35 (2H, t), 2.98-3.16 (4H, m), 3.67 (3H, s), 3.73 (3H, s), 3.91 (2H, t), 4.82 (1H, m), 6.80 (2H, d), 6.95 (2H, d).

EXAMPLE 37**2-S-(4-Carboxybutanoylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid (3-6)**

55 3-5 (0.11 g, 0.21 mmole) was treated with LiOH (0.025 g, 1.05 mmole) as described for compound 3-4 to give the desired diacid (0.105 g).

^1H NMR (300 MHz, CD_3OD) δ 1.30-1.55 (16H, m), 1.70-1.82 (4H, m), 2.20 (4H, m), 2.85 (1H, m), 3.03 (2H, m), 3.13 (1H, dd), 3.30 (1H, m), 3.92 (2H, m), 4.62 (1H, m), 6.81 (2H, d), 7.12 (2H, d).

This diacid (0.105 g) was dissolved in 30 mL EtOAc and treated with HCl gas as described for compound 2-2. The resulting solid was purified by flash chromatography on silica gel eluting with 90:8:8 ethanol/ $\text{NH}_4\text{OH}/\text{H}_2\text{O}$ to provide pure 3-6 as a white solid.

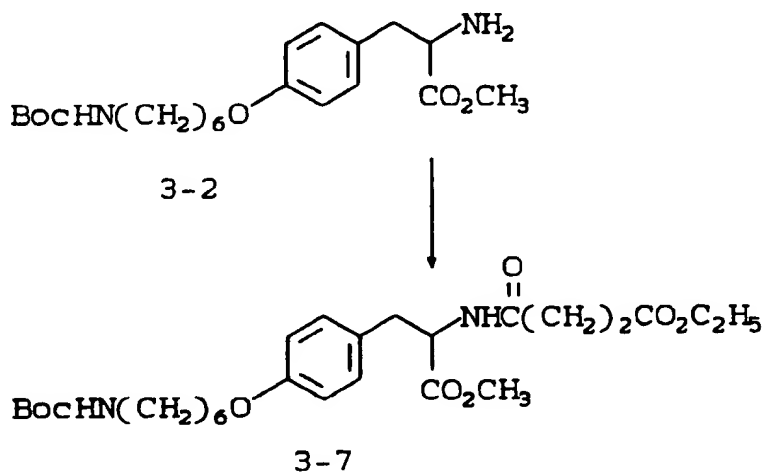
^1H NMR (300 MHz, CD_3OD) δ 1.42 (2H, m), 1.50 (2H, m), 1.63 (2H, m), 1.76 (4H, m), 2.17 (4H, m), 2.85 (3H, m), 3.16 (1H, m), 4.0 (2H, t), 4.48 (1H, m), 6.78 (2H, d), 7.12 (2H, d).

Analysis for $\text{C}_{20}\text{H}_{30}\text{N}_2\text{O}_6 \cdot 1.2 \text{ H}_2\text{O}$

Calc.: C=57.73, H=7.85, N=6.73

Found: C=57.66, H=7.21, N=6.83.

EXAMPLE 38

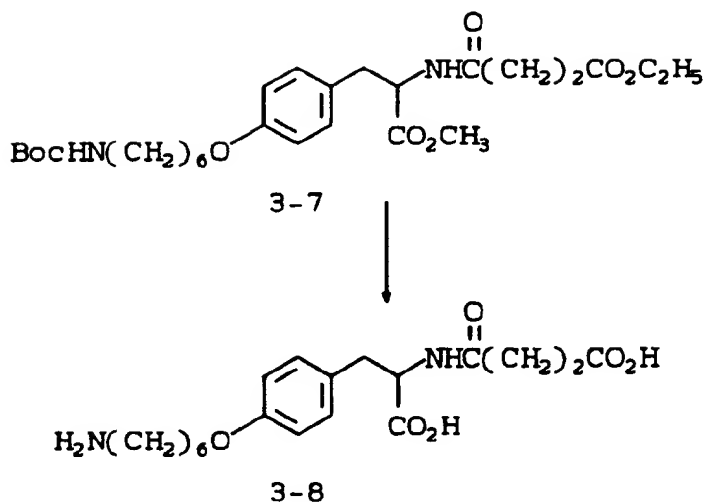


Methyl 2-S-(3-Carboethoxypropanoyl)amino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionate (3-7)

3-2 (0.562 g, 1.42 mmole) was dissolved in 15 mL EtOAc and treated with NaHCO_3 (0.36 g, 4.27 mmole) and 3-carboethoxypropanoyl chloride (0.235 g, 1.42 mmole) with stirring overnight. The reaction mixture was diluted with 150 mL EtOAc and the organic phase was washed with H_2O , brine and dried (Na_2SO_4). Solvent removal gave a residue that was purified by flash chromatography on silica gel eluting with 98:2 $\text{CHCl}_3/\text{CH}_3\text{OH}$ to give pure 3-7 (0.5 g).

^1H NMR (300 MHz, CDCl_3) δ 1.26 (3H, t), 1.35-1.61 (16H, m), 1.76 (2H, m), 2.48 (2H, m), 2.63 (2H, m), 3.05 (2H, m), 3.11 (2H, m), 3.72 (3H, s), 3.92 (2H, t), 4.13 (2H, q), 4.82 (2H, m), 6.80 (2H, d), 7.00 (2H, d).

EXAMPLE 39

2-S-(3-Carboxypropanoyl)amino-3-[4-(6-aminohexyloxy)-phenyl]propionic acid hydrochloride (3-8)

3-7 (0.58 g, 1.11 mmole) was treated with LiOH as described for 3-3 to give 2-S-(carboxypropanoyl)amino-3-[4-(6-N-t-butyloxycarbonylaminohexyloxyphenyl)]propionic acid (0.44 g) as a foam.

¹H NMR (300 MHz, CD₃OD) δ 1.32-1.58 (16H, m), 1.77 (2H, m), 2.40 (4H, m), 2.89 (1H, m), 3.0-3.16 (3H, m), 3.33 (1H, m), 3.90 (2H, t), 4.42 (1H, m), 6.78 (2H, d), 7.11 (2H, d).

This acid (0.435 g) was treated with HCl gas in EtOAc (30 mL) as described for 2-2 to give a foam that was triturated with EtOAc to give pure 3-8 (0.25 g) as a white solid.

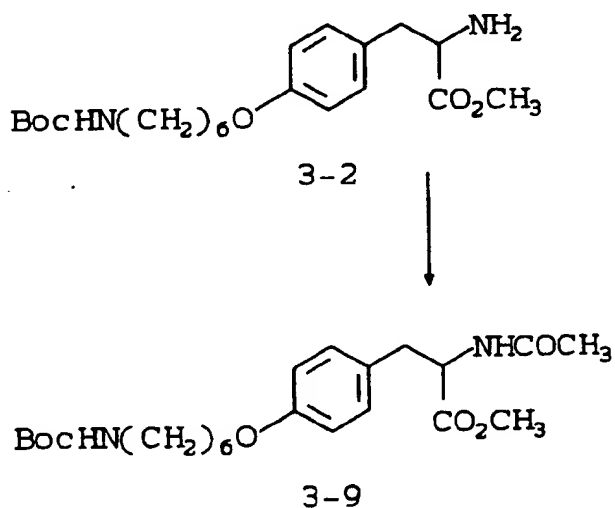
¹H NMR (300 MHz, CD₃OD) δ 1.4-1.6 (4H, m), 1.76 (2H, m), 2.46 (4H, m), 2.92 (3H, m), 3.14 (1H, m), 3.30 (1H, m), 3.96 (2H, m), 4.60 (1H, m), 6.81 (2H, d), 7.14 (2H, d).

Analysis for C₁₉H₂₈N₂O₅·HCl·0.5 H₂O

Calc.: C=53.58, H=7.10, N=6.58

Found: C=53.18, H=6.93, N=6.27.

EXAMPLE 40

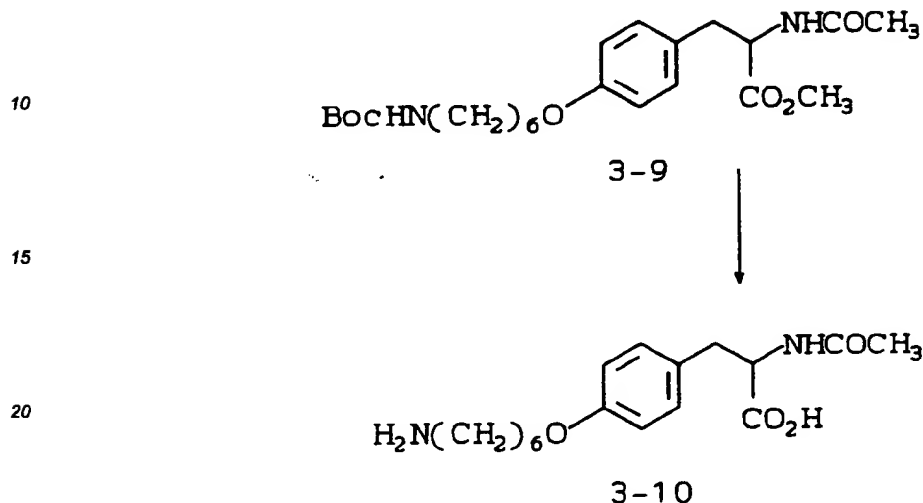
Methyl 2-S-(Acetylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionate (3-9)

3-2 (0.562 g, 1.42 mmole) was treated with acetyl chloride (0.112 g, 4.27 mmole) as described for 3-7 to give a yellow oil. This was purified by flash chromatography on silica gel eluting with 98:2 CHCl₃/CH₃OH to

give pure 3-9 (0.58 g) as a clear oil.

^1H NMR (300 MHz, CDCl_3) δ 1.30-1.56 (14H, m), 1.78 (2H, m), 2.00 (3H, s), 3.05-3.16 (4H, m), 3.73 (3H, s), 3.92 (2H, t), 4.84 (1H, m), 6.80 (2H, d), 6.98 (2H, d).

5 EXAMPLE 41



25 2-S-(Acetylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid hydrochloride (3-10)

3-9 (0.58 g, 1.33 mmole) was treated with LiOH (0.16 g, 6.64 mmole) as described for 3-3 to give 2-S-(acetylamino)-3-[4-(6-N-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid (0.485 g) as a white solid.

30 ^1H NMR (300 MHz, CD_3OD) δ 1.35-1.53 (16H, m), 1.75 (2H, m), 1.90 (3H, s), 2.86 (1H, m), 3.00-3.15 (3H, m), 3.30 (1H, m), 3.93 (2H, t), 4.59 (1H, m), 6.82 (2H, d), 7.12 (2H, d).

This compound (0.485 g) was dissolved in 30 mL EtOAc and treated with HCl gas as described for 2-2 to give a residue that was triturated with EtOAc to provide pure 3-10 (0.4 g) as a white solid.

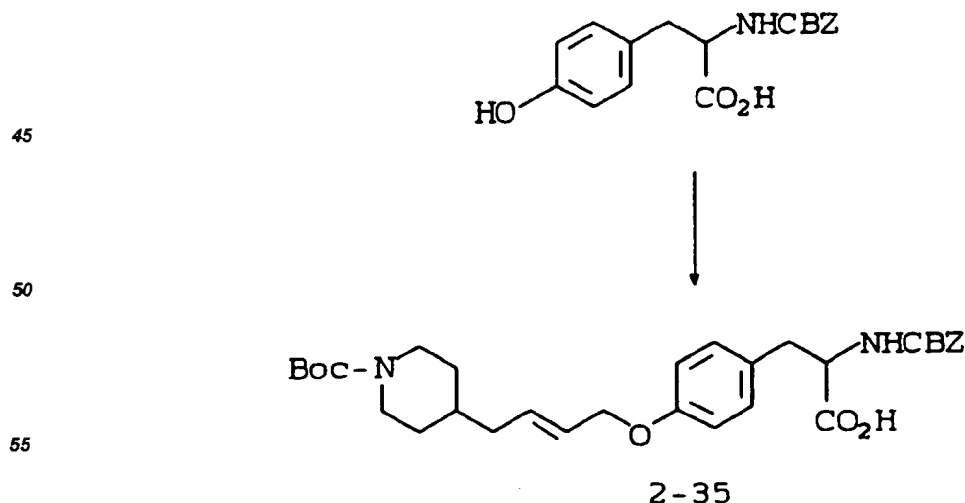
^1H NMR (300 MHz, CD_3OD) δ 1.42-1.60 (4H, m), 1.66 (2H, m), 1.70 (2H, m), 1.90 (3H, s), 2.82 (1H, m), 2.92 (2H, m), 3.12 (1H, dd), 3.30 (1H, m), 3.95 (2H, t), 4.60 (1H, m), 6.82 (2H, d), 7.13 (2H, d).

35 Analysis for $\text{C}_{17}\text{H}_{28}\text{N}_2\text{O}_4 \cdot \text{HCl} \cdot \text{H}_2\text{O}$

Calc.: C=54.17, H=7.76, N=7.43

Found: C=54.30, H=7.71, N=7.09.

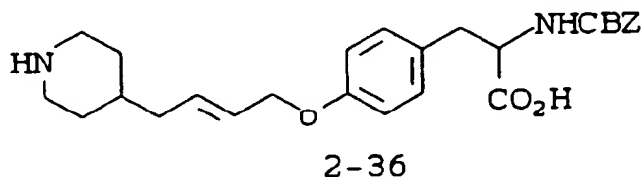
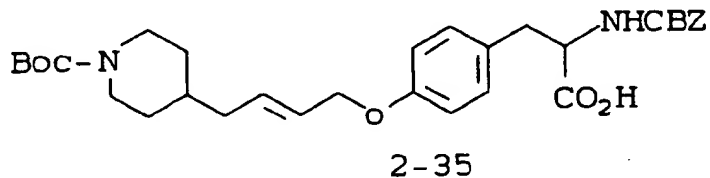
40 EXAMPLE 42



2-S-(Benzyloxycarbonylamino)-3-[4-(4-N-t-butyloxycarbonylpiperidin-4-yl)but-2-enyloxyphenyl]propionic acid (2-35)

N-CBZ-L-tyrosine (0.48 g, 0.0014 mmole) was alkylated with (4-N-t-butyloxycarbonylpiperidin-4-yl)-but-2-enyl bromide (0.424 g, 1.35 mmole) as described for 2-1. Crude product was purified by flash chromatography on silica gel eluting with 97:3:1 CHCl₃/CH₃OH/HOAc to give pure 2-35 as an oil.

¹H NMR (300 MHz, CDCl₃) δ 1.00-1.21 (4H, m), 1.40-1.55 (14H, m), 2.00-2.15 (2H, m), 2.61-2.75 (2H, m), 4.02-4.14 (3H, m), 4.57 (2H, m), 4.63 (1H, m), 5.15 (2H, m), 5.32 (1H, m), 5.58 (1H, m), 5.62-5.70 (2H, m), 6.72 (2H, t), 7.00 (2H, d).

EXAMPLE 432-S-(N-Benzyloxycarbonylamino)-3-[4-(4-piperidin-4-yl)-but-2-enyloxyphenyl]propionic acid (2-36)

2-35 (0.5 g) was dissolved in 25 mL EtOAc and treated with HCl gas as described for 2-15 to provide a residue that was titrated with ether to give 2-36. A small sample was purified by HPLC to give 2-36 as the trifluoroacetate salt.

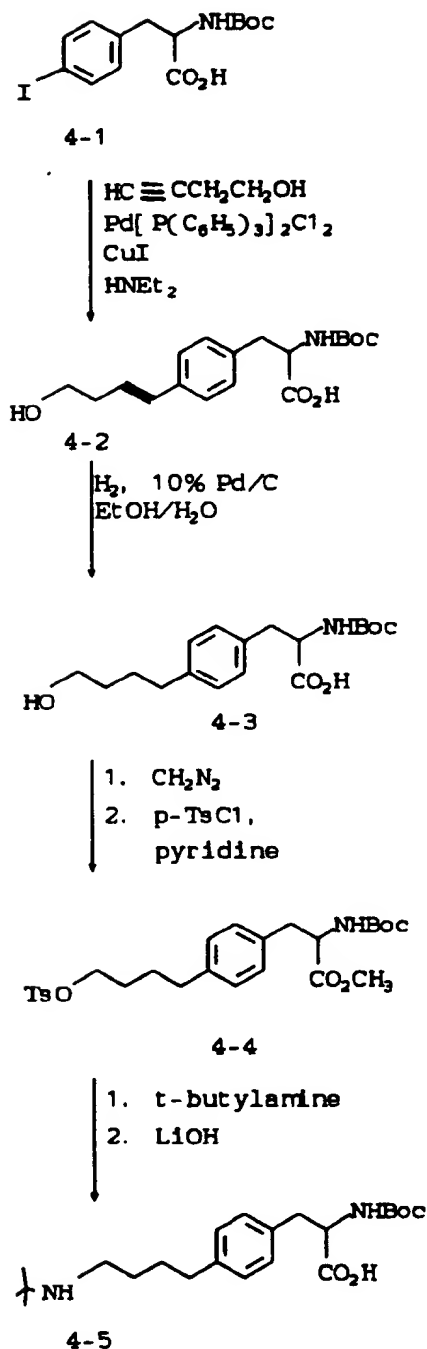
¹H NMR (300 MHz, D₂O) 7.2 (2H, m), 7.1 (4H, m), 6.7 (2H, d), 5.5 (2H, m), 5.1 (1H, d), 4.8 (1H, d), 4.2 (3H, bs), 3.2 (1H, d), 2.8 (3H, m), 2.25 (2H, 6t), 1.8 (2H, m), 1.4 (3H, m), 1.2 (1H, m), 0.9 (2H, m).

Analysis for C₂₈H₃₂N₂O₅

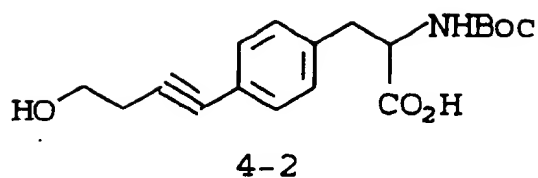
Calc.: C=57.87, H=5.68, N=4.75

Found: C=57.98, H=5.79, N=4.61

SCHEME 4

2-S-(N-t-Butyloxycarbonylamino)-3-[4-(4-hydroxybut-1-ynyl)phenyl]propionic acid (4-2)

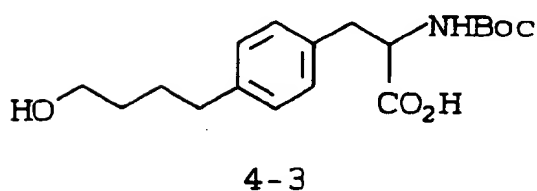
N-BOC-4-iodo-L-phenylalanine (4-1) (1.0 g, 2.55 mmole) was dissolved in diethylamine under N_2 and treated with 3-butyne-1-ol (0.23 mL, 3.06 mmole), $[\text{Pd}(\text{P}(\text{C}_6\text{H}_5)_3)_2\text{Cl}_2]$ (0.089 g, 0.127 mmole) and CuI (0.012 g, 0.064 mmole). After 3 hours the solvent was evaporated, the residue dissolved in water (pH = 11) and extract d
 55 with ethyl acetate. The water layer was then acidified to pH 3, extracted with ethyl acetate. This organic extract was dried and vaporated to give 0.8 g crude 4-2. $R_f = 0.47$ in 97/3/1 $\text{CHCl}_3/\text{CH}_3\text{OH}/\text{HOAc}$, ninhydrin stain. ^1H NMR (300 MHz, CDCl_3) δ 7.35 (2H, d), 7.1 (2H, d), 6.4 (1H, broad), 5.0 (1H, d), 4.6 (1H, m), 3.8 (2H, t), 3.1 (2H, m), 2.65 (2H, t), 1.4 (9H, s).

EXAMPLE 47

10

H₂, 10% Pd/C
Et OH/H₂O

15

2-S-(N-t-Butyloxycarbonylamino)-3-[4-(4-hydroxybutyl)phenyl]propionic acid (4-3)

30

4-2 (0.40 g, 1.2 mmole) was dissolved in an ethanol/water solution (25 mL) and was treated with 10% Pd/C (0.1 g) and H₂ on a Parr apparatus. After 2 hours the solution was filtered and evaporated. Column chromatography on silica gel (94:5:1 CHCl₃/CH₃OH/HOAc) yielded 0.321 g (80%) of 4-3. R_f=0.57 in 97:3:1 CHCl₃/CH₃OH/HOAc ninhydrin stain.

35

¹H NMR (300 MHz, CDCl₃) δ 7.1 (s, 4H), 4.95 (1H, m), 4.9 (1H, broad), 4.55 (1H, m), 3.65 (2H, t), 3.1 (2H, m), 1.6 (4H, m), 1.4 (9H, s).

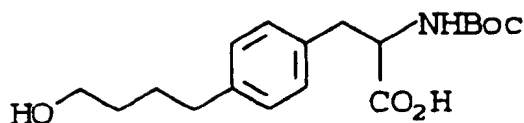
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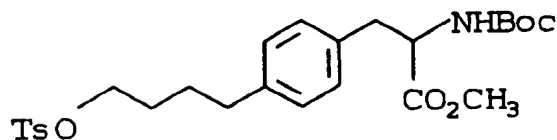
55

EXAMPLE 48



4-3

1. CH_2N_2
2. p-TsCl,
pyridine



4-4

Methyl 2-S-(N-t-Butyloxycarbonylamino)-3-[4-(4-tosyloxybutyl)phenyl]propionate (4-4)

4-3 (0.285 g, 0.85 mmole) was dissolved in CH_2Cl_2 (10 mL) cooled to 0°C , and treated with CH_2N_2 solution. After 10 minutes the reaction was quenched with MgSO_4 , filtered and evaporated to provide ester used in the next reaction. $R_f=0.5$ in 92:8:1 $\text{CHCl}_3/\text{CH}_3\text{OH}/\text{HOAc}$, ninhydrin stain.

^1H NMR (300 MHz, CDCl_3) δ 7.05 (d, $J=7.8$ Hz, 2H), 7.0 (d, $J=7.8$ Hz, 2H), 5.0 (1H, m), 4.55 (1H, m), 3.69 (3H, s), 3.6 (2H, $J=6.2$ Hz, t), 3.0 (2H, m), 2.6 (2H, $J=7.5$ Hz, t), 1.7 (4H, m), 1.4 (9H, s).

This ester was dissolved in 10 mL CH_2Cl_2 and added at -78°C to a solution prepared from treating p-toluenesulfonyl chloride (0.14 g, 0.67 mmole) in CH_2Cl_2 at -78°C with pyridine (0.1 mL, 1.35 mmole) for 10 minutes. The reaction was allowed to warm to room temperature over 1.0 hour and then water was added. The organic layer was separated, dried, and evaporated. Column chromatography 97:3:1 on silica gel eluting with $\text{CHCl}_3/\text{CH}_3\text{OH}/\text{HOAc}$ gave 4-4 (0.27 g, 70%). $R_f=0.85$ 97:3:1 $\text{CHCl}_3/\text{CH}_3\text{OH}/\text{HOAc}$.

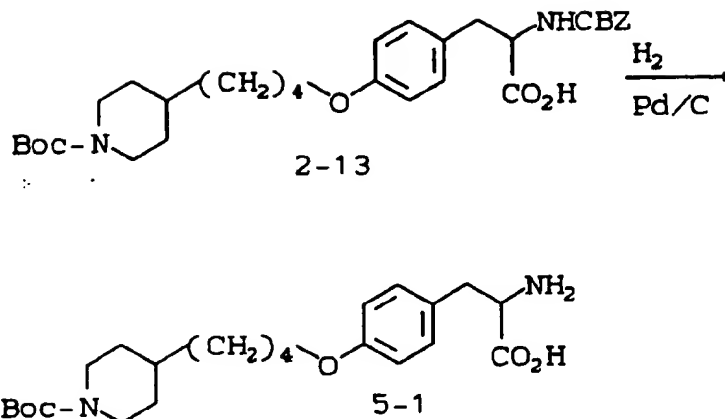
^1H NMR (300 MHz, CDCl_3) δ 7.88 (2H, $J=7.2$ Hz, d), 7.74 (2H, $J=7.2$ Hz, d), 7.38 (2H, $J=\text{Hz}$, d), 7.30 (2H, $J=8$ Hz, d), 5.0 (1H, m), 4.5 (1H, m), 4.0 (2H, $J=5.3$ Hz, t), 3.67 (3H, s), 3.0 (2H, m), 2.5 (2H, t), 2.0 (3H, s), 1.6 (4H, m), 1.4 (9H, s).



Found: C=56.73, H=7.51, N=5.58.

51

EXAMPLE 50

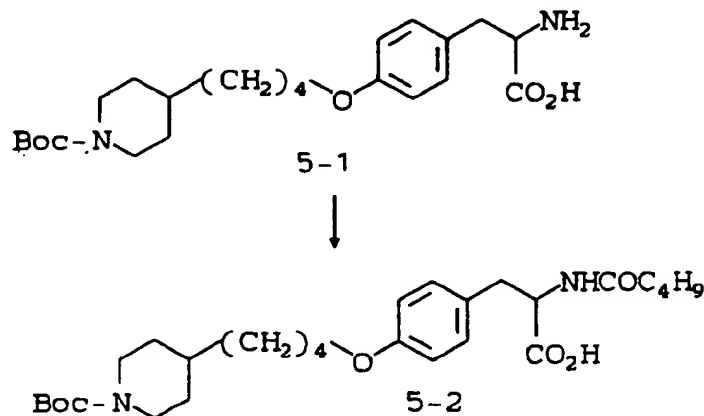
2-S-Amino-3-[4-(4-N-t-butyloxycarbonylpiperidin-4-yl)butyloxyphenyl]propionic acid (5-1)

20

2-13 (2.0 g) was dissolved in 100 mL EtOH, and 0.2 g 10% Pd/C was charged. This suspension was hydrogenated at balloon pressure overnight. Solvent removal provided 5-1 (1.36 g) as a white solid.

¹H NMR (300 MHz, CD₃OD), δ 0.97-1.12 (2H, m), 1.20-1.54 (14H, m), 1.72 (4H, m), 2.71 (2H, m), 2.90-3.00 (1H, m), 3.22 (1H, dd), 3.30 (1H, m), 3.71 (1H, m), 3.95-4.10 (4H, m), 6.88 (2H, d), 7.21 (2H, d).

EXAMPLE 51

2-S-(Pentanoylamino)-3-[4-(4-N-t-butyloxycarbonylpiperidin-4-yl)butyloxyphenyl]propionic acid (5-2)

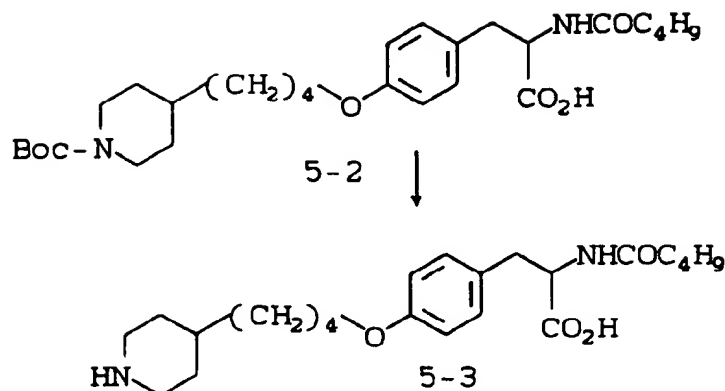
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5-1 (1.05 g, 2.5 mmole) was added to a cold solution of 1 N NaOH (2.5 mL) in 20 mL H₂O and stirred at 0-10 degrees C for 5 minutes to give a clear solution. Then, pentanoyl chloride (0.332 g, 2.75 mmole) was added dropwise followed by NaHCO₃ (0.231 g, 2.75 mmole) and the resulting mixture was stirred vigorously at 0-10° C for 1 hour. The reaction mixture was diluted with H₂O (75 mL), acidified to pH 2-3 with 10% KHSO₄ and extracted with EtOAc. This extract was filtered, washed with brine, dried (Na₂SO₄) and the solvent removed to give an oil. This was purified by flash chromatography on silica gel eluting with 97:3:1 CHCl₃/CH₃OH/HOAc to give pure 5-2 (0.44 g) as a clear oil.

50

¹H NMR (300 MHz, CD₃OD) δ 0.90 (3H, t), 1.20-1.62 (16H, m), 1.72 (2H, m), 2.14 (2H, m), 2.30 (8H, m), 2.65-2.90 (4H, m), 3.30 (1H, m), 3.93 (2H, m), 4.61 (1H, m), 6.81 (2H, d), 7.12 (2H, d).

EXAMPLE 52

2-S-(Pentanoylamino)-3-[4-(4-piperidin-4-yl)butyloxy]-phenyl]propionic acid hydrochloride (5-3)

5-2 (0.449 g), was dissolved in 30 mL EtOAc and treated with HCl gas at -10° C as described for 2-2. The resulting solid was triturated with 40 mL Et₂O to give pure 5-3 (0.36 g) as a white solid.

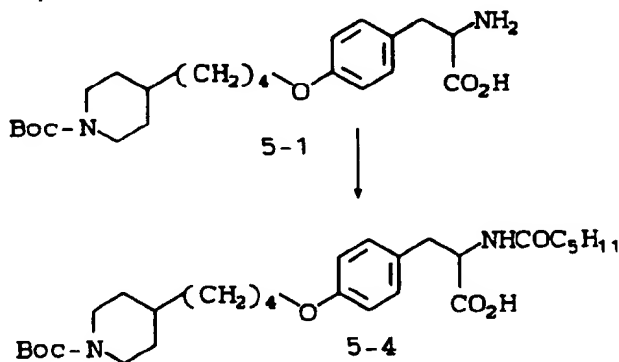
¹H NMR (300 MHz, CD₃OD) δ 0.85 (3H, t), 1.19 (2H, m), 1.30-1.65 (9H, m), 1.73 (2H, m), 1.95 (2H, m), 2.15 (2H, m), 2.80-3.02 (3H, m), 3.14 (1H, dd), 3.30-3.40 (3H, m), 3.95 (2H, t), 4.61 (1H, m), 6.82 (2H, d), 7.13 (2H, d).

Analysis for C₂₃H₃₈N₂O₄·HCl·0.75 H₂O

Calc.: C = 60.77, H = 8.54, N = 6.16

Found: C = 60.97, H = 8.39, N = 6.06.

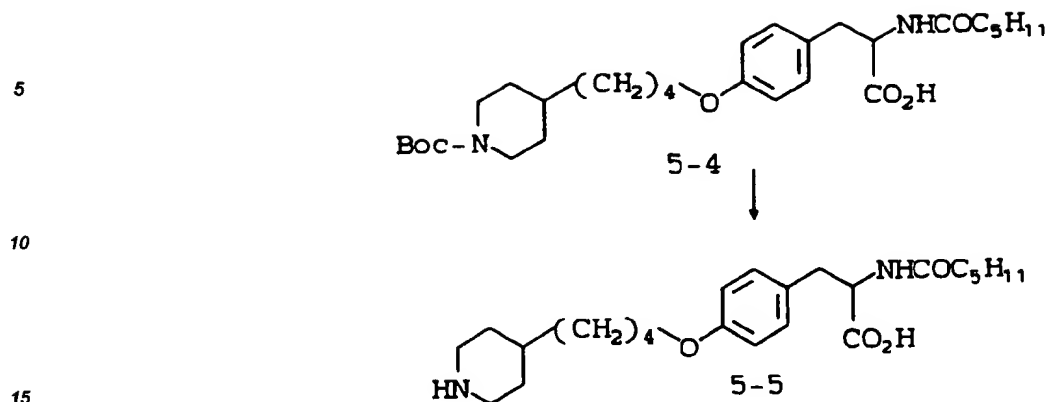
EXAMPLE 53

2-S-(Hexanoylamino)-3-[4-(4-N-t-butyloxycarbonylpiperidin-4-yl)butyloxyphenyl]propionic acid (5-4)

5-1 (0.41 g) was treated with hexanoyl chloride (0.21 mL, 1.50 mmole) as described for 5-2. Crude product was purified by flash chromatography on silica gel eluting with 97:3:1 CHCl₃/CH₃OH/HOAc to give pure 5-4 (0.20 g).

¹H NMR (300 MHz, CD₃OD) δ 0.85 (3H, t), 0.97-1.35 (8H, M), 1.37-1.53 (12H, m), 1.60-1.80 (4H, m), 2.13 (2H, t), 2.80 (2H, m), 2.85 (1H, m), 3.12 (1H, dd), 3.90 (2H, t), 4.04 (2H, d), 4.62 (1H, m), 6.80 (2H, d), 7.12 (2H, d).

EXAMPLE 54



2-S-(Hexanoylamino)-3-[4-(4-piperidin-4-ylbutoxy)phenyl]propionic acid (5-5)

20 5-4 (0.199 g) was dissolved in 25 mL EtOAc and treated with HCl gas as described for compound 2-2 to provide pure 5-5 (48 mg).

¹H NMR (300 MHz, CD₃OD) δ 0.84 (3H, t), 1.08-1.20 (4H, m), 1.35 (4H, m), 1.52 (4H, m), 1.77 (2H, m), 1.92 (2H, d), 2.16 (2H, t), 2.80-3.2 (3H, m), 3.15 (1H, dd), 3.40-3.52 (2H, m), 3.92 (2H, t), 4.61 (1H, m), 6.81 (2H, d), 7.13 (2H, d).

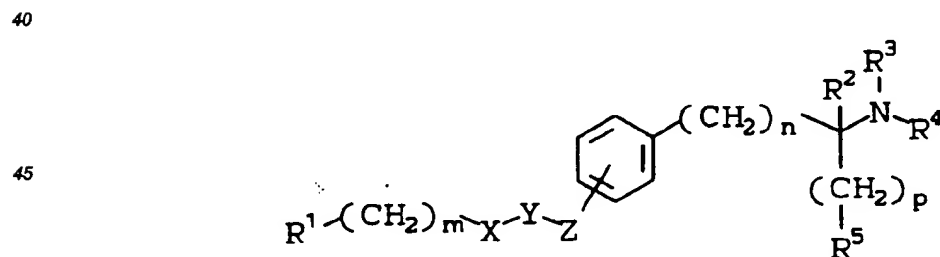
25 Analysis for $C_{28}H_{39}N_2O_8F_3 \cdot 0.55 H_2O \cdot 0.30 TFA$

Calc.: C = 55.39, H = 7.06, N = 4.86

Found: C = 55.38, H = 7.03, N = 4.85.

Sample alternative protecting groups that can be used in the preparation of the present invention include benzyl ester, cyclohexyl ester, 4-nitrobenzyl ester, t-butyl ester, 4-pyridylmethyl ester, benzyloxycarbonyl, isonicotinylloxycarbonyl, O-chlorobenzyloxycarbonyl, p-nitrobenzyloxycarbonyl, p-methoxybenzyloxycarbonyl, t-butoxycarbonyl, t-amylloxycarbonyl, isobornylloxycarbonyl, adamantylloxycarbonyl, 2-(4-biphenyl)-2-propyloxycarbonyl and 9-fluorenylmethoxycarbonyl.

In addition to those compounds specifically exemplified above, additional compounds of the present invention are set forth in tabular form below. These compounds are synthesized by use of the synthetic routes and methods described in the above Schemes and Examples and variations thereof well known to those of ordinary skill in the art, and not requiring undue experimentation. All variables listed in the Tables below are with reference to the following generic structure:



Example	R ¹	R ²	R ³	R ⁴	R ⁵	X	Y	Z	m	n	p
55		H	H	-CH ₂ C ₆ H ₅	CO ₂ H	CH ₂	0	0	3	1	1
56		H	CH ₃	COCH ₃	CO ₂ H	CH ₂	0	-	3	1	1
57		CH ₂ OCH ₃	CH ₂ C ₆ H ₅	COC ₂ H ₅	CO ₂ H ₅	CH ₂	CH ₂	CH ₂	4	2	1
58		(CH ₂) ₂ C ₆ H ₅		COCH ₂ CH(CH ₃) ₂	COCH ₃	CH ₂	CH ₂	-	5	3	1
59		H	(CH ₂) ₂ OCH ₂ CH ₃	(CH ₂) ₂ C ₆ H ₅	$\begin{matrix} \text{C-OH} \\ \\ \text{S} \end{matrix}$	CH ₂	0	CH ₂	2	2	2
60			H	CO(CH ₂) ₂ OCH ₃	$\begin{matrix} \text{C-O-C-O-C(CH}_3)_3 \\ \quad \\ \text{O} \quad \text{O} \end{matrix}$	CH ₂	CH ₂	-	3	1	1

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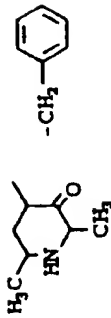



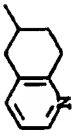

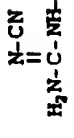

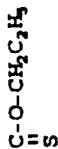

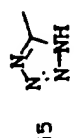



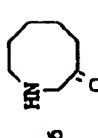



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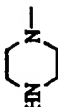
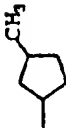

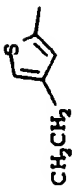

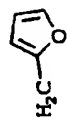
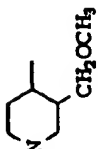

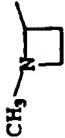

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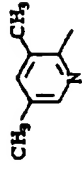

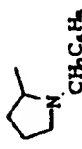
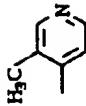

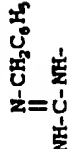


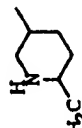


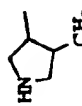
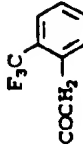
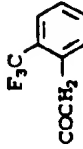
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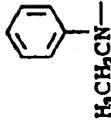
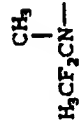

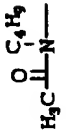







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Example	R ¹	R ²	R ³	R ⁴	R ⁵	X	Y	Z	m	n	P
61			C ₆ H ₅	COO(CH ₂) ₂ C ₆ H ₅	CO ₂ CH(CH ₃) ₂	CH ₃	O	-	2	3	1
62		CH ₂ CH ₂ CF ₃	(CH ₂) ₂ SO ₂ CH ₃	CH ₂ SO ₂ CH ₂ C ₂ H ₅		CHOH	O	-	3	1	1
63		H	(CH ₂) ₂ CN	H	CONHCH ₂ CO ₂ CH ₃	NH		-	0	6	2
64		C ₆ H ₄	(CH ₂) ₂ NO ₂			NCH ₃		-	2	0	10
65			CF ₂ CF ₃			O	CH ₂	-	3	1	1
66			(CH ₂) ₂ NHCH ₃			SO ₂	CH=CH	-	1	2	3

Example	R ¹	R ²	R ³	R ⁴	R ⁵	X	Y	Z	m	n	p
67			$(\text{CH}_2)_4\text{SCH}_2\text{CH}_3$	$\text{COCH}_2\text{C}_6\text{H}_5$	$\text{O}=\text{P}(\text{OH})$	CH_2	$\text{C}=\text{S}$	CH_2	4	2	1
68			$(\text{CH}_2)_4$ 	H_2C 	$\text{O}=\text{P}(\text{OH})_2$	$\text{C}=\text{O}$	0	-	3	3	0
69		$\text{CH}_2\text{SC}_6\text{H}_5$	$(\text{CH}_2)_3\text{CO}_2\text{CH}_3$	COC_3H_7	$-\text{C}-\text{O}-\text{C}-\text{O}-\text{CH}_3$ \parallel O	$\text{C}=\text{S}$	NHCH_3	-	4	1	3
70	$\text{C}_6\text{H}_5\text{CH}_2\text{NHC}-$ \parallel NH	$\text{CH}_2\text{CH}_2\text{SO}_2\text{CH}_3$	$(\text{CH}_2)_2\text{CO}_2\text{H}$		$-\text{C}-\text{O}-\text{C}_2\text{H}_5$ \parallel O	CH_2	$\text{C}=\text{S}$	CH_2	3	4	1
71		$\text{CH}_2\text{CH}_2\text{NHCH}_3$	$(\text{CH}_2)_3\text{NH}(\text{CH}_2)_2\text{OCH}_3$	$\text{COOC}_{10}\text{H}_{21}$		CH_2	SO_2	-	2	2	2

Example	R ¹	R ²	R ³	R ⁴	R ⁵	X	Y	Z	m	n	p
72		CH ₃		H	CO ₂ H	CH ₂	NCOCH ₃	-	1	1	1
73		C ₂ H ₅		(CH ₂) ₂ CO ₂ H		O	C=O	0	3	2	1
74		C ₃ H ₇			COOC ₂ H ₅	CH ₂	SO ₂	CH ₂	1	0	4
75		C ₄ H ₉	C ₃ H ₇			CH ₂	CH ₂	0	3	1	3
76		CH ₂ C ₆ H ₅	C ₁₀ H ₂₁			CH=CH	CH ₂	-	2	2	2

Example	R ¹	R ²	R ³	R ⁴	R ⁵	X	Y	Z	m	n	p
77		(CH ₂) ₃ C ₆ H ₅	(CH ₂) ₃ -SO ₂ -C ₆ H ₅	CH ₂ (CH ₂) ₂ CO ₂ CH ₃	-COOC ₆ H ₅	CH ₂	-C-	-	1	2	4
78		(CH ₂) ₂ C ₆ H ₅	CH ₂ CH ₂ F	-C-(CH ₂) ₂ OCH ₃		-C=C-	CH ₂	0	4	2	1
79		(CH ₂) ₃ -NHCOOH	CH ₃		-CO ₂ CH ₂ C ₆ H ₅		CH ₂	-	3	2	0
80	C ₆ H ₅ (CH ₂) ₂ NH-	(CH ₂) ₃ SC ₆ H ₅	C ₂ H ₅			CH ₂	CH ₂	-	0	2	3
81		CH ₂ - 	(CH ₂) ₂ CH ₂ NO ₂	CH ₃		CH ₂	CH ₂	-	1	1	1

The test procedures employed to measure the anti-platelet aggregating activity of the compounds of the present invention are described below.

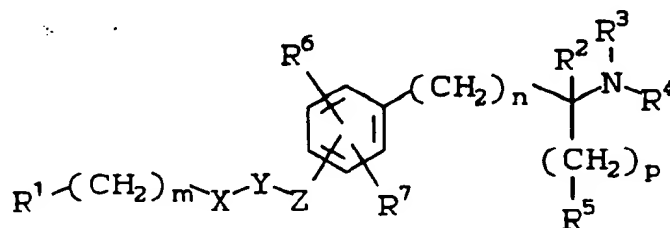
EXAMPLE 82

Blood was drawn into 0.1 volumes of acid-citrate-dextrose (85 mM sodium citrate, 64 mM citric acid, 110 mM dextrose) by venipuncture from normal human volunteers. Platelet-rich plasma was prepared by centrifugation at 400 x g for 12 minutes. PGE1 (5 mg/ml) was added and platelets were collected by centrifugation at 800 x g for 12 minutes. The platelet pellet was resuspended into human platelet buffer (140 mM NaCl, 7.9 mM KCl, 3.3 mM Na₂HPO₄, 6 mM HEPES, 2% bovine serum albumin, 0.1 % dextrose, pH 7.2) and filtered over Sepharose 2B that was previously equilibrated in human platelet buffer. Platelets were counted and adjusted to 2 x 10⁸/ml with human platelet buffer. Human fibrinogen (10-100 mg/ml and CaCl₂ (1 mM) were added and aggregation was initiated by the addition of 10 mM ADP. Aggregation was monitored by the initial rate of increase of light transmittance.

While the invention has been described and illustrated in reference to certain preferred embodiments thereof, those skilled in the art will appreciate that various changes, modifications and substitutions can be made therein without departing from the spirit and scope of the invention. For example, effective dosages other than the preferred doses as set forth hereinabove may be applicable as a consequence of variations in the responsiveness of the mammal being treated for severity of clotting disorders or emboli, or for other indications for the compounds of the invention indicated above. Likewise, the specific pharmacological responses observed may vary according to and depending upon the particular active compound selected or whether there are present pharmaceutical carriers, as well as the type of formulation and mode of administration employed, and such expected variations or differences in the results are contemplated in accordance with the objects and practices of the present invention. It is intended, therefore, that the invention be limited only by the scope of the claims which follow and that such claims be interpreted as broadly as is reasonable.

Claims

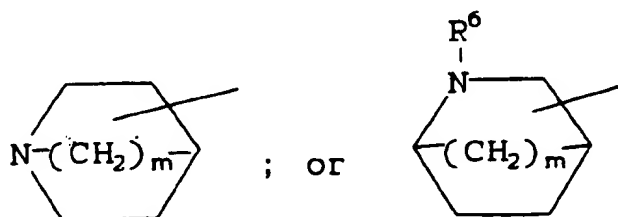
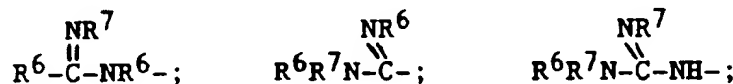
1. A compound of the formula



and the pharmaceutically acceptable salts thereof, wherein

R¹ is

a four to eight member heterocyclic ring containing 1, 2, 3 or 4 hetero atoms wherein said heteroatoms are N, O or S and wherein said heterocyclic ring is optionally substituted at any atom by H, R⁶ or R⁷;



NR⁶R⁷ wherein R⁶ and R⁷ are independently
hydrogen,

C₁₋₁₀ alkoxy carbonyl or unsubstituted or substituted C₁₋₁₀ alkyl and cycloalkyl wherein said substituents
are

- 5 C₁₋₁₀ alkoxy,
- C₁₋₁₀ alkoxyalkyl,
- C₁₋₁₀ alkoxyalkyloxy,
- C₁₋₁₀ alkoxy carbonyl,
- 10 C₁₋₁₀ alkyl carbonyl,
- C₀₋₈ alkylaminocarbonyl,
- C₁₋₁₀ aralkyl carbonyl,
- C₁₋₁₀ alkylthiocarbonyl,
- C₄₋₁₀ aralkylthiocarbonyl,
- 15 thiocarbonyl,
- C₁₋₁₀ alkoxythiocarbonyl,
- aryl,
- 5 to 6 membered saturated heterocyclic rings of 1, 2, 3 or 4 hetero atoms wherein said hetero atoms
are taken from the group consisting of N, O and S,
- 20 C₁₋₄ alkanoylamino,
- C₁₋₆ alkoxy carbonyl-C₀₋₈ alkylamino,
- C₁₋₁₀ alkylsulfonylamino,
- C₄₋₁₀ aralkylsulfonylamino,
- 25 C₄₋₁₀ aralkyl,
- C₁₋₁₀ alkaryl,
- C₁₋₁₀ alkylthio,
- C₄₋₁₀ aralkylthio,
- C₁₋₁₀ alkylsulfinyl,
- 30 C₄₋₁₀ aralkylsulfinyl,
- C₁₋₁₀ alkylsulfonyl,
- C₄₋₁₀ aralkylsulfonyl,
- aminosulfonyl,
- C₁₋₁₀ alkylaminosulfonyl,
- 35 C₄₋₁₀ aralkylsulfonylamino,
- oxo,
- thio,
- unsubstituted or mono- or
di-substituted 1-ethenyl, 2-ethenyl or 3-propenyl wherein said substituents are selected from the
40 group consisting of hydrogen, C₁₋₁₀ alkyl and C₇₋₁₀
- aralkyl,
- carboxy,
- hydroxy,
- amino,
- 45 C₁₋₆ alkylamino,
- C₁₋₆ dialkylamino,
- halogen, where halogen is defined as Cl, F, Br, or I,
- nitro, or
- cyano,

50 and further wherein said N can additionally be substituted to form a quaternary ammonium ion wherein
said substituent is as previously defined for R⁶ and R⁷;

R² and R³ are independently

hydrogen,

aryl or

55 unsubstituted or substituted C₀₋₁₀ alkyl or cycloalkyl wherein said substituent is

C₁₋₁₀ alkoxyalkyl,

aryl,

a 4 to 8 membered heterocyclic ring containing 1, 2, 3 or 4 hetero atoms, wherein said hetero atoms are taken from the group consisting of N, O and S,

5 C₄₋₁₀ aralkyl,
C₁₋₁₀ alkaryl,
carboxy,
C₁₋₁₀ alkylcarbonyl,
C₁₋₁₀ alkylthiocarbonyl,
10 C₄₋₁₀ aralkylcarbonyl,
C₄₋₁₀ aralkylthiocarbonyl,
C₁₋₆ alkoxycarbonyl,
C₄₋₁₀ aralkoxycarbonyl,
C₁₋₆ alkoxy,
C₄₋₁₀ aralkoxy,
15 C₁₋₆ alkylamino,
C₁₋₁₂ dialkylamino,
C₁₋₆ alkanoylamino,
C₄₋₁₂ aralkanoylamino,
C₄₋₁₀ aralkylamino;

20 R⁴ is

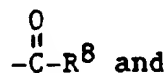
hydrogen,
aryl,
C₁₋₁₀ alkyl or cycloalkyl
25 C₄₋₁₀ aralkyl,
arylcarbonyl, aminocarbonyl,
C₁₋₁₀ alkylcarbonyl, C₁₋₆ alkylaminocarbonyl,
C₁₋₁₀ alkylthiocarbonyl, C₁₋₆ dialkylaminocarbonyl,
C₁₋₁₀ alkoxythiocarbonyl, arylC₁₋₆ alkylaminocarbonyl,
30 C₁₋₁₀ alkoxycarbonyl,
C₄₋₁₀ aralkylcarbonyl,
C₄₋₁₀ aralkoxycarbonyl,
C₁₋₁₀ carboxylalkyl and

35 further wherein any of the substituents for R⁴ may be substituted by one or more substituents selected from the group as defined for R⁶, or an L- or D-amino acid joined by an amide linkage;

R⁵ is

a four to eight membered saturated or unsaturated heterocyclic ring containing 1, 2, 3 or 4 hetero atoms wherein said hetero atoms are N, O, or S or,

40



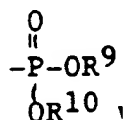
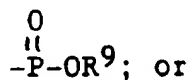
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wherein R⁸ is

50 hydroxy,
C₁₋₁₀ alkyloxy,
C₁₋₁₀ alkaryloxy,
C₄₋₁₀ aralkyloxy,
C₄₋₁₀ aralkylcarbonyloxy,
55 C₁₋₁₀ alkoxyalkyloxy,
C₁₋₁₀ alkoxyalkylcarbonyloxy,
C₁₋₁₀ alkoxycarbonyloxyalkyl,

C₁₋₁₀ alkylcarbonyloxyalkyloxy,
 an L- or D-amino acid joined by an amide linkage, and wherein the carboxylic acid moiety of said
 amino acid is as the free acid or is esterified by C₁₋₆alkyl.



wherein R⁹ and R¹⁰ are selected from the group consisting of
 hydrogen, C₁₋₁₀ alkyl and C₄₋₁₀ aralkyl;

X and Y are optional substituents that, when present, are

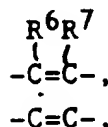
NR⁶,

O,

S,

SO,

SO₂,

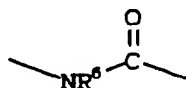
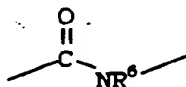


oxo,

aryl,

thiono,

unsubstituted or substituted C₁₋₁₅ alkyl or cycloalkyl wherein said substituents are independently
 R⁶ and R⁷,



-NR⁶-SO₂-, -SO₂-NR⁶-, or

a 4- to 8- membered heterocyclic ring containing 1, 2, 3, or 4 heteroatoms wherein said atoms are N, O,
 or S and wherein said ring is independently substituted at any atom with R⁶;

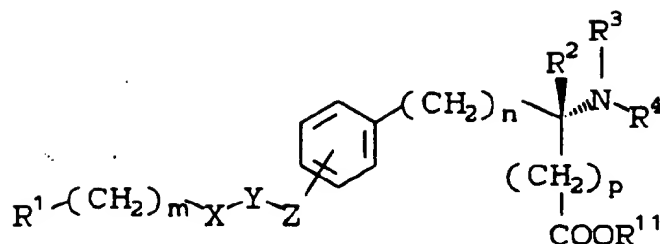
Z is an optional substituent that, when present, is independently chosen as defined by X and Y;

m is an integer of from zero to ten;

n is an integer of from zero to ten; and

p is an integer of from zero to three.

2. A compound of the structural formula



and the pharmaceutically acceptable salts thereof, wherein

R¹ is

a four to eight member heterocyclic ring containing 1, 2, 3 or 4 hetero atoms wherein said heteroatoms are N, O, or S and wherein said heterocyclic ring is optionally substituted by hydrogen, C₁₋₁₀ alkyl;

or

NR⁶R⁷ wherein R⁶ and R⁷ are independently

hydrogen,

C₁₋₁₀ alkoxy carbonyl or

unsubstituted or substituted C₁₋₁₀ alkyl wherein said substituent is

C₁₋₁₀ alkoxy,

C₁₋₁₀ alkoxy carbonyl,

aryl,

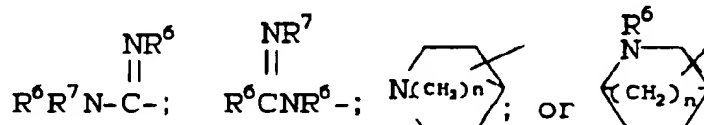
C₄₋₁₀ aralkyl,

C₁₋₁₀ alkaryl,

carboxy,

hydroxy or

amino,



and further wherein said N can additionally be substituted to form a quaternary ammonium ion;

R^2 and R^3 are independently

hydrogen or

C₁₋₁₀ alkyl; or

C₄₋₁₀ aralkyl;

R⁴ is

hydrogen,

C₁₋₁₀ alkyl,

C₄₋₁₀ aralkyl,

arylcarbonyl,

aralkylcarbonyl

C₁₋₁₀ alkylcarbo

C₁₋₁₀ alkoxycarbonyl

C₄₋₁₀ aralkylcarbonyl.

C₄₋₁₀ aralkoxycarbonyl

and further wherein any of the substituents for R⁴ may be substituted by one or more substituents

from the group defined as R⁶ in Claim 1;

R¹¹ is

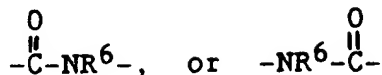
hydrogen or

C₁₋₁₀ alkyl;

X and Y are independently

O.

S, SO
 SO₂,
 aryl,
 -CH=CH-,
 OXO,

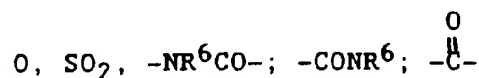


-NR⁶SO₂-; -SO₂NR⁶-

unsubstituted or substituted C₁₋₁₅ straight or branched alkyl either substituted or unsubstituted with

carboxy,
 hydroxy,
 C₁₋₁₀ alkoxy, or

a 4- to 6- membered heterocyclic ring containing 1, 2 or 3 heteroatoms chosen from N, O or S,
 Z is an optional substituent that, when present, is



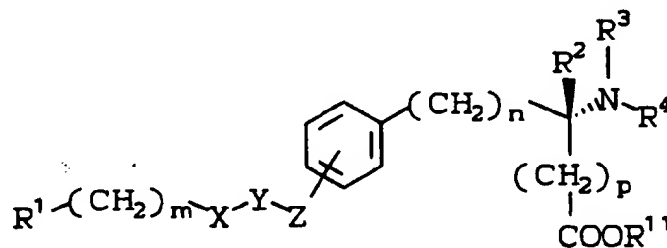
C₁₋₁₀ straight or branched alkyl;

m is an integer of from zero to six;

n is an integer of from zero to six; and

p is an integer of from zero to three.

3. A compound of the structural formula



and the pharmaceutically acceptable salts thereof, wherein

R¹ is

a five to six member heterocyclic ring containing 1 or 2 heteroatoms wherein said heteroatoms are N and wherein said heterocyclic ring is optionally substituted by C₁₋₅ alkyl; or NR⁶R⁷ wherein R⁶ and R⁷ are independently

hydrogen,

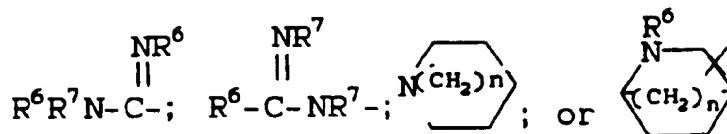
unsubstituted or substituted C₁₋₁₀ alkyl

wherein said substituent is

C₁₋₁₀ alkoxy, carbonyl,

aryl,

C₁₋₁₀ aralkyl,



and further wherein said N can additionally be substituted to form a quaternary ammonium ion;

R² and R³ are hydrogen;

R⁴ is

arylcarbonyl,

C₁₋₁₀ alkylcarbonyl,

aralkylcarbonyl

C₁₋₁₀ alkoxycarbonyl,

C₄₋₁₀ aralkylcarbonyl,

C₄₋₁₀ aralkoxycarbonyl or

and further wherein the substituents for R⁴ may be unsubstituted by one or more substituents from the group defined as R⁶ in Claim 1;

R¹¹ is

hydrogen or

C₁₋₁₀ alkyl;

X and Y are independently

O, SO₂, aryl; NR⁶CO-; -CONR⁶-

-CH=CH-,

unsubstituted or substituted C₁₋₁₅ straight or branched alkyl wherein said substituent is hydroxy, or

a 4- to 6- membered heterocyclic ring containing 1 or 2 heteroatoms chosen from N, O or S;

Z is an optional substituent that, when present, is

O or

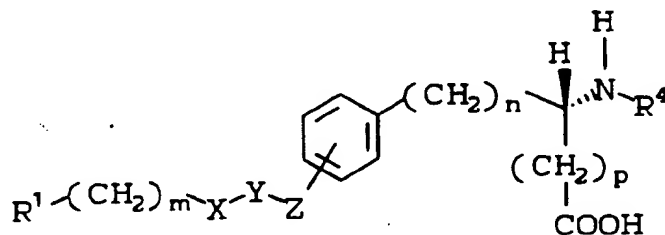
C₁₋₁₀ straight or branched alkyl;

m is an integer of from zero to six;

n is an integer of from zero to one; and

p is an integer of from zero to one.

4. A compound of the structural formula



and the pharmaceutically acceptable salts thereof, wherein

R¹ is

a six member saturated heterocyclic ring containing 1 or 2 heteroatoms wherein said heterocyclic atoms are N and wherein said heterocyclic ring is optionally substituted by C₁₋₅ alkyl; or

NR⁶R⁷ wherein R⁶ and R⁷ are independently hydrogen or

C₁₋₁₀ alkyl;

R⁴ is

arylcarbonyl,

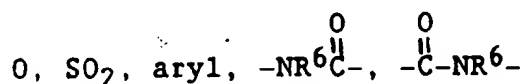
C₁₋₁₀ alkylcarbonyl,

C₄₋₁₀ aralkylcarbonyl

C₁₋₁₀ alkoxycarbonyl,

C₄₋₁₀ aralkylcarbonyl or

C_{4-10} aralkoxycarbonyl,
and further wherein any of the substituents for R^4 may be substituted by one or more substituents
from the group defined as R^6 in Claim 1,
X and Y are independently



-CH=CH- or
 C_{1-10} straight or branched alkyl;
Z is an optional substituent that, when present, is
O or
 C_{1-5} straight or branched alkyl;
m is an integer of from zero to six;
n is one; and
p is zero.

5. A compound as claimed in Claim 1, selected from the group consisting of:

2-S-(6-N-Benzyloxycarbonylamino)-3-[4-(3-chloropropoxy)phenyl] propionic acid;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(N,N,2,2-tetramethyl-1,3-propanediamino)propyloxyphenyl]propionic acid;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(3-N-pyrrolidinypropyloxy)phenyl]propionic acid ;

2-S-(N-Benzyloxycarbonylamino)-[4-(3-N-methyl-N-benzylaminopropoxyphenyl)]propionic acid;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(4-piperazinyl)butyloxyphenyl]propionic acid;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(1,1,4,4-tetramethylbutylamino)propyloxyphenyl]propionic acid;

2-S-(N-Benzyloxycarbonyl)-3-[4-(4-methylpiperazin-1-yl)propyloxyphenyl]propanoic acid;

2-(N-Benzyloxycarbonylamino)-3-[4-(5-bromopentyloxy)phenyl]propionic acid ;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(4-piperazin-1-yl)pentyloxyphenyl]propionic acid;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(6-aminohexyloxyphenyl)]propionic acid hydrochloride;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(7-aminohexyloxy)phenyl]propionic acid hydrochloride;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(8-aminooctyloxy)phenyl]propionic acid;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(5-aminopentyloxy)phenyl]propionic acid hydrochloride;

2-S-(N-Benzyloxycarbonylamino)-3-[4-(4-piperidinylbutyloxy)phenyl]propionic acid;

2-S-Phenylcarbonylamino-3-[4-(6-aminohexyloxy)phenyl]propionic acid hydrochloride;

2-S-Phenethylcarbonylamino-3-[4-(6-aminohexyloxy)phenyl]propanoic acid hydrochloride;

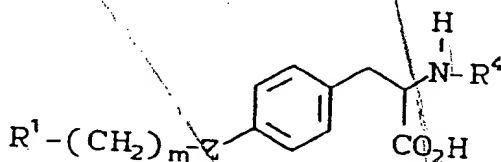
2-S-(Phenylacetyl)amino-3-[4-(6-aminohexyloxy)phenyl]propionic acid;

2-S-(2-Carboxy-3-phenylpropionylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid;

- 2-S-(Hexanoylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid Hydrochloride;
- 2-S-(Naphthanoylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid;
- 5 2-S-(Butanoylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid;
- 2-S-(Heptanoylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid hydrochloride ;
- 2-S-(5-Phenylpentanoylamino)-3-[4-(6-t-butyloxycarbonylaminohexyloxy)phenyl]propionic acid;
- 10 2-S-(5-Phenylpentanoylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid hydrochloride;
- 2-S-(3-Carboxypropanoyl)amino-3-[4-(6-aminohexyloxy)phenyl]propionic acid hydrochloride;
- 15 2-S-(Acetylamino)-3-[4-(6-aminohexyloxy)phenyl]propionic acid hydrochloride;
- 2-S-(N-Benzoyloxycarbonylamino)-3-[4-(4-piperidinyl)but-2-enyloxyphenyl]propionic acid;
- 2-S-(N-t-Butyloxycarbonylamino)-3-[4-(4-hydroxybut-1-ynyl)phenyl]propionic acid;
- 20 2-S-(N-t-Butyloxycarbonylamino)-3-[4-(4-hydroxybutyl)phenyl]propionic acid;
- 2-S-(N-t-Butyloxycarbonylamino)-3-[4-(4-t-butylaminobutyl)phenyl]propionic acid;
- 25 2-S-(Pentanoylamino)-3-[4-(4-piperidin-4-ylbutyloxy)phenyl]propionic acid hydrochloride;
- 2-S-(Hexanoylamino)-3-[4-(4-piperidin-4-ylbutyloxy)phenyl]propionic acid;
- 2-S-(5-Aminopentanoyl)amino-3-[4-(6-aminohexyloxy)phenyl]propionic acid dihydrochloride;
- 30 Methyl 2-S-(4-Carbomethoxybutanoyl)amino-3-[4-(N-t-butyloxycarbonylaminohexyloxy)phenyl]propionate;
- 2-S-(4-Carboxybutanoylamino)-3-[4-(6-aminohexyloxy) phenyl]propionic acid; and
- 35 2-S-(3-Carboxypropanoyl)amino-3-[4-(6-aminohexyloxy) phenyl]propionic acid hydrochloride.
6. The use of a compound as claimed in claim 1 for the manufacture of a medicament for blocking fibrinogen from acting at its receptor site in a mammal.
- 40 7. The use of a compound as claimed in claim 1 for the manufacture of a medicament for prevention or treatment of thrombus and embolus formation.
8. The use of a compound as claimed in claim 1 for the manufacture of a medicament for inhibiting aggregation of blood platelets.
- 45 9. The use of a compound as claimed in claim 1 together with an anti-coagulant agent for the manufacture of a medicament for prevention or treatment of thrombus and embolus formation.
10. The use of a compound as claimed in claim 1 together with an anti-coagulant agent for the manufacture of a medicament for inhibiting aggregation of blood platelets.
- 50 11. The use of a compound as claimed in claim 1 together with a thrombolytic agent for the manufacture of a medicament for prevention or treatment of thrombus and embolus formation.
- 55 12. The use of a compound as claimed in claim 1 together with a thrombolytic agent for the manufacture of a medicament for inhibiting aggregation of blood platelets.
13. The use of a compound as claimed in claim 1 together with a platelet anti-aggregation agent for the manufacture of a medicament for inhibiting aggregation of blood platelets.

ufacture of a medicament for prevention or treatment of thrombus and embolus formation.

14. The use of a compound as claimed in claim 1 together with a platelet anti-aggregation agent for the manufacture of a medicament for inhibiting aggregation of blood platelets.
15. A pharmaceutical composition, comprising a compound as claimed in Claim 1, and a pharmaceutically acceptable carrier.
16. A pharmaceutical composition comprising the compound as claimed in Claim 1, a pharmaceutically acceptable carrier and a compound taken from the group consisting of thrombolytic agents, platelet anti-aggregation agents and anti-coagulant agents.
17. The composition as claimed in Claim 15 or Claim 16, in which said pharmaceutically acceptable carrier consists of a sustained release pharmaceutical formulation.
18. The compounds of Claim 1 for use in inhibiting the binding of fibrinogen to blood platelets, inhibiting the aggregation of blood platelets, treating thrombus formation or embolus formation, or preventing thrombus or embolus formation in a mammal.
19. The compounds of Claim 5 for use in inhibiting the binding of fibrinogen to blood platelets, inhibiting the aggregation of blood platelets, treating of thrombus formation or embolus formation, or preventing thrombus or embolus formation in a mammal.
20. A compound as claimed in Claim 1 of formula



wherein

R¹ is

a five or 6-membered heterocyclic ring wherein said heteroatom is N and wherein said heterocyclic ring is optionally substituted by hydrogen or C₁-₆ alkyl, or

NR⁶R⁷ wherein R⁶ and R⁷ are independently hydrogen, C₁-₁₀ alkyl or C₄-₁₀ arylalkyl;

R⁴ is

arylcarbonyl,

C₁-₁₀ alkylcarbonyl,

C₁-₁₀ alkoxy carbonyl,

C₄-₁₀ aralkylcarbonyl,

C₄-₁₀ aralkoxy carbonyl wherein R⁴ is unsubstituted or substituted with R⁶ as previously defined;

Z is chosen from:

O, -NR⁶CO-, -CONR⁶-, or CH₂; and

m is an integer of from one to six



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 91 30 8793

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
X	EP-A-0 071 926 (FUJISAWA PHARMACEUTICAL CO. LTD.) * page 51 - page 54 *	1-4	C07C271/22 A61K31/40 C07D295/088 C07C271/16
X	ARZNEIMITTEL FORSCHUNG. vol. 39-I, no. 3, March 1989, AULENDORF DE pages 328 - 334; I. SETNIKAR ET AL: 'METABOLISM AND EXCRETION OF 14-C-TIROPRAIDE AFTER SINGLE INTRAVENOUS OR PERORAL ADMINISTRATION TO THE RAT' *SUMMARY; COMPOUNDS CR1809 AND CR1931*	1-4, 6-20	C07C233/87 C07C233/51 C07D211/22 A61K31/325 A61K31/445
X	ARZNEIMITTEL FORSCHUNG. vol. 38-II, no. 12, December 1988, AULENDORF DE pages 1815 - 1819; I. SETNIKAR: 'TIROPRAIDE AND METABOLITES IN BLOOD AND PLASMA AFTER INTRAVENOUS OR PERORAL ADMINISTRATION OF 14-C-TIROPRAIDE TO THE RAT' *SUMMARY; COMPOUNDS CR1809 AND CR1931*	1-4, 6-20	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C07C C07D
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 20 DECEMBER 1991	Examiner SANCHEZ Y GARCIA J.
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 1503 (01.92) (P0401)

